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# INTERIM REPORT February 25, 1993

**FOR** 

## **BIOVENTING FIELD INITIATIVE**

AT

## HANSCOM AIR FORCE BASE, MASSACHUSETTS

to

Captain Catherine M. Vogel
Department of the Air Force
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by

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## TABLE OF CONTENTS

1.0	INTROL	DUCTIO	N	
			ves	
			scriptions	
			Building 1639 Site	-
		1.2.2	Building 1812 Site	-
				-
2.0	BUILDI	NG 1639	)	•
	2.1	Chronol	logy of Events and Site Activities	ì
			Groundwater Measurements	ì
		2.1.2	Soil Gas Survey	
		2.1.3	Vent Well, Monitoring Point, and Thermocouple Installation	
		2.1.3		10
		215		10
		2.1.5		
	. 22	Poculte		10
	2.2	2 2 1	Soil and Soil Cos Analyses	12
		2.2.1		12
		2.2.2	Soil Gas Permeability and Radius of Influence	12
		2.2.3	In Situ Respiration Test	12
		2.2.4	Bioventing Demonstration	16
3.0	מות חווא	NG 1812	• • • • • • • • • • • • • • • • • • • •	1 C
5.0	3 1	Chronol		18
	3.1			18
		3.1.1	Soil Gas Survey	18
		3.1.2		18
		3.1.3		20
		3.1.7		22
	3.2	Peculte :		22
	3.2	2 2 1	Soil and Soil Gos Analyses	22
		3.2.1		22
		2 2 2		24 24
		3.2.3	Dioventing Demonstration	24
4 0	BACKGI	ROUND	AREA ACTIVITIES	24
	2.101101	COULD	AMAZI 11011VIIII.D	۷4
5.0	FUTURE	E WORK	· · · · · · · · · · · · · · · · · · ·	27
6.0	REFERE	NCE .	• • • • • • • • • • • • • • • • • • • •	30
APP:	ENDIX A	: TEST	PLAN FOR HANSCOM AFB, MASSACHUSETTS A	-1
A DD	ENIDIS O	. ABTAT	LAMICAL DEDONE FOR DAMA PARA COMPANY	
APP.	ENDIX E	S: ANAI	LYTICAL REPORT FOR BUILDING 1639, BUILDING 1812, AND	_
		BACK	KGROUND AREA B	-1
A DD	ENIDIV C	1. Drift.	DINC 1620 COIL CAR DEDICE ADDITION DATE	
APP.	ENDIY (	. DUIL	DING 1639 SOIL GAS PERMEABILITY DATA	-1

APPENDI	IX D: BUILDING 1639 IN SITU RESPIRATION TEST DATA	D-1
APPENDI	IX E: BUILDING 1812 SOIL GAS PERMEABILITY DATA	E-1
	LIST OF TABLES	
Table 1. Table 2. Table 3. Table 4. Table 5.	Initial Soil Gas Composition at Building 1639	7 13 14 14
Table 6. Table 7. Table 8. Table 9. Table 10.	Respiration Test at Building 1639  Initial Soil Gas Composition at Building 1812  Results From Soil Analyses for BTEX and TPH at Building 1812  Results From Soil Chemistry Analysis at Building 1812  Results of Hyperventilate <sup>™</sup> Soil Gas Permeability Analysis  Results From Soil Chemistry Analysis at Background Area	16 19 23 23 25 28
	LIST OF FIGURES	
Figure 1. Figure 2.	Schematic Diagram of Hanscom AFB	3
Figure 3.	Schematic Diagram of Building 1812 at Hanscom AFB (GS - Soil Gas Survey Point; MP - Monitoring Point)	5
Figure 4.	Cross Section of Vent Well and Monitoring Points at Building 1639 Showing Site Lithology and Construction Detail (not to scale)	9
Figure 5. Figure 6.	Radius of Influence at Building 1639	15
Figure 7.	Respiration Test at Monitoring Point H1-MPB-5.0'	17
Figure 8. Figure 9.	Site Lithology and Construction Detail (not to scale)	21 26
<i>5</i>	Respiration Test at the Background Area	29

## **INTERIM REPORT**

#### **BIOVENTING FIELD INITIATIVE**

## HANSCOM AIR FORCE BASE, MASSACHUSETTS

## 1.0 INTRODUCTION

This report describes the activities conducted at Hanscom Air Force Base (AFB), Massachusetts, as part of the Bioventing Field Initiative for the U.S. Air Force Center for Environmental Excellence (AFCEE) and the Environmental Quality Directorate of the Air Force Armstrong Laboratory. This report summarizes the results from the first phase of the study at Hanscom AFB. First-phase activities include a soil gas survey, air permeability test, in situ respiration tests, and installation of bioventing systems. The specific objectives of this Bioventing Field Initiative are described in the following section. Each site at the base is discussed individually, followed by a description of site activities at the background area.

## 1.1 Objectives

The purpose of this Bioventing Field Initiative is to measure the soil gas permeability and microbial activity at a contaminated site in order to evaluate the potential application of bioventing technology to remediate the site. The specific test objectives are stated below.

- A small-scale soil gas survey will be conducted to identify an appropriate location for installation of the bioventing system. Soil gas from the candidate site should exhibit high total petroleum hydrocarbon (TPH) concentrations, relatively low oxygen concentrations, and relatively high carbon dioxide concentrations. An uncontaminated background location also will be identified.
- The soil gas permeability of the soil and the air vent (well) radius of influence will be determined. To measure these parameters, air will be withdrawn or injected for approximately 8 hours at vent wells located in contaminated soils. Pressure changes will be monitored in an array of monitoring points.
- Immediately following the soil gas permeability test, an in situ respiration test will be conducted. Air will be injected into selected monitoring points to

aerate the soils. The in situ oxygen utilization and carbon dioxide production rates will be measured.

The data from the soil gas permeability and in situ respiration tests will be used to determine an air injection/withdrawal rate for the bioventing test. A blower will be selected, installed, and operated for 6 to 12 months, and periodic measurements of the soil gas composition will be made to evaluate the long-term effectiveness of bioventing.

## 1.2 Site Descriptions

Hanscom AFB is located in Bedford, Massachusetts. A schematic diagram of the base is shown in Figure 1. The dashed lines on the map indicate the direction from the main gate to each test site. Site H1 and Site H2 refer to Building 1639 and Building 1812, respectively. Groundwater at Hanscom AFB generally is encountered at 3 to 8 feet. The sites chosen for the bioventing test initiative are Building 1639 and Building 1812. A brief description of the sites is provided in Sections 1.2.1 and 1.2.2. A detailed description of the test sites is provided in the Test Plan in Appendix A.

#### 1.2.1 Building 1639 Site

Building 1639 is the base fuel service station (Figure 2). There have been three reported petroleum releases (gasoline, #2 fuel oil, and waste lubricating oil) at this facility since November 1990. Soil contamination at Building 1639 ranges from 599 ppm to 4,400 ppm TPH.

## 1.2.2 Building 1812 Site

The site at Building 1812 was the location of a heating oil fuel tank (Figure 3). Leakage occurred from the tank, resulting in soil contamination at this site. The tank was replaced recently, and soil TPH concentrations as high as 2,830 mg/kg have been reported.

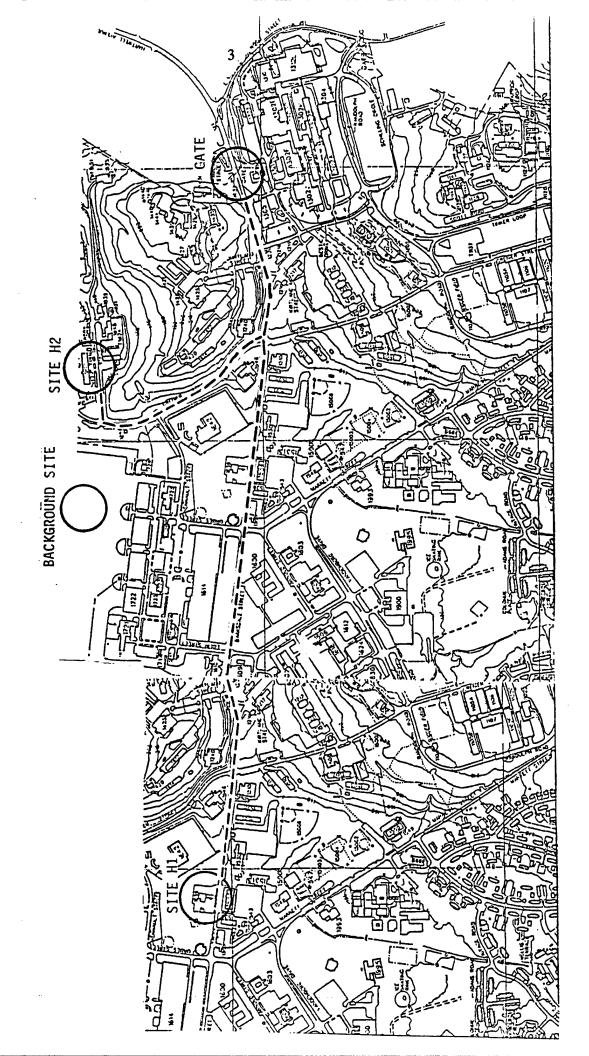


Figure 1. Schematic Diagram of Hanscom AFB

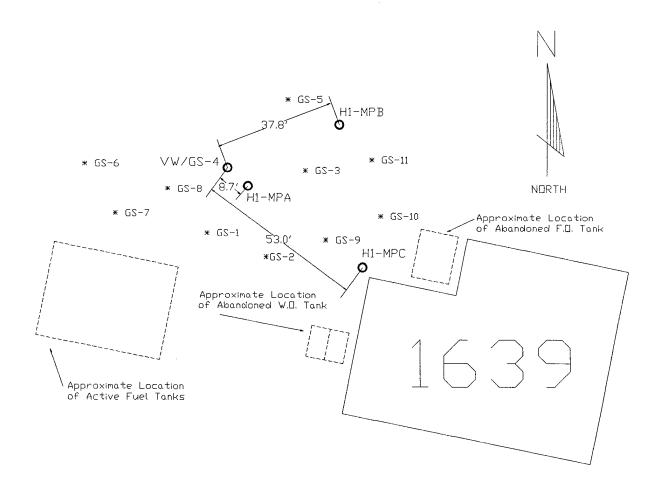


Figure 2. Schematic Diagram of Building 1639 at Hanscom AFB (GS - Soil Gas Survey Point; MP - Monitoring Point)

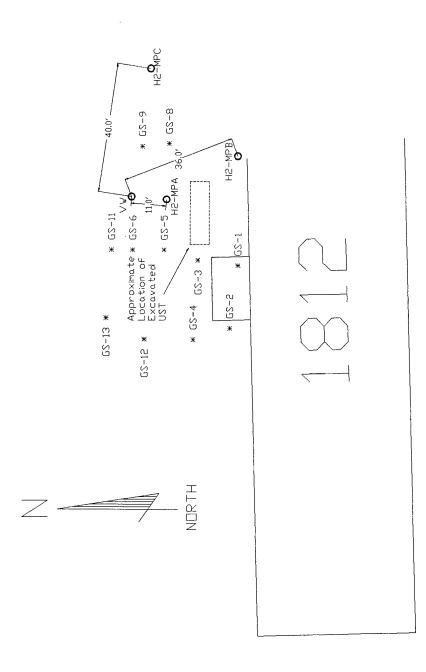


Figure 3. Schematic Diagram of Building 1812 at Hanscom AFB (GS - Soil Gas Survey Point; MP - Monitoring Point)

## 2.0 BUILDING 1639

## 2.1 Chronology of Events and Site Activities

#### 2.1.1 Groundwater Measurements

One groundwater monitoring well (12BO3) was present at Building 1639. Groundwater level was measured at this well on September 30, 1992 and was recorded at 5.75 feet.

## 2.1.2 Soil Gas Survey

A suitable site for the bioventing demonstration should have soil gas characteristics of high TPH, low oxygen, and high carbon dioxide concentrations. This composition of soil gas would indicate that oxygen-limiting conditions for microbial activity are present and that the introduction of air may enhance biodegradation of TPH.

On September 30, 1992, a limited soil gas survey was conducted at Building 1639. Soil gases were sampled by driving a %-inch-diameter stainless steel probe into the soil with a hammer drill. Soil gas was withdrawn with a vacuum pump and analyzed for oxygen, carbon dioxide, and TPH.

Measurements of oxygen and carbon dioxide in the soil gas were made with a GasTech Model 32520X with oxygen and carbon dioxide ranges of 0 to 25%. The analyzer was calibrated daily against atmospheric oxygen, atmospheric carbon dioxide, a 10% oxygen calibration standard, and a 5% carbon dioxide calibration standard. TPH was measured with a GasTech Trace Techtor with TPH ranges from 0 to 100, 0 to 1,000, and 0 to 10,000 ppm. The GasTech Trace Techtor was calibrated daily against a 4,200 ppm hexane standard.

The soil gas probes were driven to depths ranging from 2.5 to 5.0 feet at several locations at Building 1639. Table 1 provides the initial concentrations of oxygen, carbon dioxide, and TPH for the various locations at Building 1639. Oxygen concentrations varied from 0.8 to 21%, whereas TPH concentrations ranged from 410 ppm to greater than 20,000 ppm. The oxygen concentrations in the soil gas indicate that some areas at this site are oxygen-limited and may respond to bioventing.

Table 1. Initial Soil Gas Composition at Building 1639

Soil Gas Survey (GS) Point	Depth (ft)	Oxygen (%)	Carbon Dioxide (%)	TPH (ppm)
GS-1	2.5	19.0	2.5	15,600
	5.0	20.2	0.70	6,000
GS-2	2.5	16.0¹	5.8	>20,000
	5.0	2.0	18.5	>20,000
GS-3	2.5	1.0	18.5	>20,000
	5.0	3.0	17.0	>20,000
GS-4	2.5	21.0¹	0.10	920
	5.0	1.0	23.0	>20,000
GS-5	2.5	16.2	5.8	660
	5.0	15.0	7.0	720
GS-6	2.5	20.0	1.8	410
GS-7	2.5	21.0	0.50	900
GS-8	2.5	19.5¹	1.0	5,600
	5.0	19.2¹	1.8	8,800
GS-9	2.5	4.0¹	13.5	>20,000
	5.0	0.80 <sup>1</sup>	19.5	>20,000
GS-10	2.5	13.5¹	6.8	>20,000
	5.0	0.80 <sup>1</sup>	17.0	>20,000
GS-11	2.5	16.0¹	4.3	>20,000

Pressure reading on sampling pump was high. Measured oxygen concentration may not be representative of actual soil gas oxygen concentrations. Actual oxygen concentration is likely to be lower.

## 2.1.3 Vent Well, Monitoring Point, and Thermocouple Installation

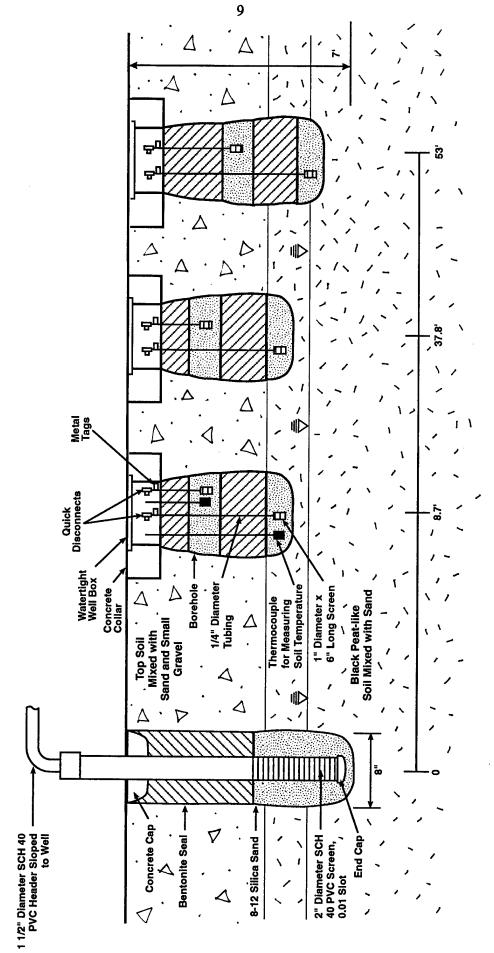
On October 3, 1992, one vent well (VW) and three monitoring points (MPs) were installed, and soil samples were collected for analyses. The monitoring points were labeled as follows: H1-MPA; H1-MPB; and H1-MPC. The locations of the vent well and monitoring points are shown in Figure 2. A cross section of the vent well and monitoring points showing site lithology and construction detail is shown in Figure 4.

The vent well was installed at a depth of 7.0 feet into an 8-inch-diameter borehole. The vent well consisted of Schedule 40 2-inch-diameter polyvinyl chloride (PVC) piping with 3.0 feet of tenslot screen. The annular space corresponding to the screened area of the well was filled with silica sand, and the annular space above the screened interval was filled with bentonite to prevent short-circuiting of air to or from the surface.

Soil gas probes consisted of ¼-inch tubing with a 1-inch-diameter, 6-inch screened area. The annular space corresponding to the screened area was filled with silica sand, whereas the interval between the screened areas was filled with bentonite, as was the annular space from the shallowest monitoring point to the ground surface. The monitoring points were installed at depths as follows:

- Monitoring point H1-MPA was installed at a depth of 5.0 feet into an 8-inch-diameter borehole. The monitoring point was screened to two depths: 2.5 and 5.0 feet.
- Monitoring point H1-MPB was installed at a depth of 5.0 feet into an 8-inch-diameter borehole. The monitoring point was screened to two depths: 2.5 and 5.0 feet.
- Monitoring point H1-MPC was installed at a depth of 6.0 feet into an 8-inch-diameter borehole. The monitoring point was screened to two depths: 3.5 and 6.0 feet.

A Type K thermocouple was installed with monitoring points H1-MPA-2.5' and H1-MPA-5.0'.



MPC

MPB

MPA

**Vent Well** 

Figure 4. Cross Section of Vent Well and Monitoring Points at Building 1639 Showing Site Lithology and Construction Detail (not to scale)

## 2.1.4 Soil and Soil Gas Sampling and Analyses

Split-spoon soil samples were collected at depths of 4.0 to 4.5 feet and 4.5 to 5.0 feet from the vent well borehole and were labeled H1-VW-4'-4.5' and H1-VW-4.5'-5.0', respectively. A soil sample also was collected from monitoring point A at a depth of 3.0 to 4.0 feet and was labeled H1-A-3'-4'. The samples were sent under chain of custody to Engineering-Science Inc., Berkeley Laboratory for analysis of benzene, toluene, ethylbenzene, and xylenes (BTEX); TPH; alkalinity; moisture content; pH; iron; total phosphorous; total Kjeldahl nitrogen; and particle size analysis.

Soil vapor samples were collected from all monitoring points and were labeled H1-A-2.5, H1-A-5, H1-B-2.5, H1-B-5, H1-C-3.5, and H1-C-6. These samples were sent under chain of custody to Air Toxics, Ltd., in Rancho Cordova, California, for analysis of BTEX and TPH.

## 2.1.5 Soil Gas Permeability and Radius of Influence

A detailed description of the method for conducting a soil gas permeability test, including equations to compute k, the soil gas permeability, is given in the Test Plan and Technical Protocol (Hinchee et al., 1992).

Prior to air injection, the monitoring points were allowed to set up for 24 hours. Air was injected with a portable 1-horsepower (HP) explosion-proof positive displacement blower unit. After air injection was initiated, pressure readings were taken approximately every 1 to 2 minutes for the first hour, then approximately every 10 minutes for the following hour. The Hyperventilate<sup>TM</sup> computer model was used to calculate the soil gas permeability.

## 2.1.6 In Situ Respiration Test

Immediately following the soil gas permeability test, air containing approximately 1% helium was injected into the soil for approximately 24 hours, beginning on October 6. Air was injected concurrently into the background monitoring well to measure the natural biodegradation of organic material in the soil. The setup for the in situ respiration test is described in the Test Plan and Technical Protocol (Hinchee et al., 1992). The pump used for air injection was a 1/3-HP diaphragm pump. Air and helium were injected through the following monitoring points at the depths indicated: H1-MPA-5.0'; H1-MPB-2.5'; H1-MPB-5.0'; and H1-MPC-6.0'. After the air/helium injection was

turned off, the respiration gases were monitored periodically. The respiration test was terminated on October 10.

Helium concentrations were measured during the in situ respiration test to quantify helium leakage to or from the surface around the monitoring points. Helium loss over time is attributed to either diffusion or leakage. A rapid drop in helium concentration followed by a leveling is an indication of leakage. A gradual loss along with an apparent first-order curve is an indicator of diffusion. As a rough estimate, the diffusion of gas molecules is inversely proportional to the square root of the molecular weight of the gas. Based on molecular weights of 4 for helium and 32 for oxygen, helium gas diffuses about 2.8 times faster than oxygen, or the diffusion of oxygen is 0.35 times the rate of helium diffusion. As a general rule, we have found that if helium concentrations are at least 50 to 60% of the initial levels at test completion, measured oxygen uptake rates are representative. Greater helium loss indicates a problem, and oxygen utilization rates are not considered representative.

To compare data from one site to another, a stoichiometric relationship of the oxidation of the hydrocarbon was assumed. Hexane was used as the representative hydrocarbon for the organic contaminant. The stoichiometric relationship is given by:

$$C_6H_{14} + 9.5O_2 - 6CO_2 + 7H_2O$$
 (1)

Based on the utilization rates (% per day), the biodegradation rates in terms of milligrams as a hexane equivalent per kilogram of soil per day were computed using the equation below by assuming a soil porosity of 0.2 and a bulk density of 1,440 kg/m<sup>3</sup>.

$$K_{\beta} = \frac{-K_o A D_o C}{100} \tag{2}$$

where:  $K_{g}$  = biodegradation rate (mg/kg/day)

K<sub>o</sub> = oxygen utilization rate (percent per day)

A = volume of air/kg of soil, in this case 300/1,440 = 0.21

 $D_o$  = density of oxygen gas (mg/L) assumed to be 1,330 mg/L

C = mass ratio of hydrocarbon to oxygen required for mineralization, assumed to be 1/3.5 from the above stoichiometric equation.

## 2.2 Results and Discussion

## 2.2.1 Soil and Soil Gas Analyses

Results of the soil analyses for BTEX and TPH at Building 1639 are presented in Table 2. The analytical report for this site is presented in Appendix B. Concentrations of the BTEX compounds in soil samples ranged from 0.015 mg/kg (ethylbenzene) up to 12 mg/kg (total xylenes), whereas TPH concentrations ranged from below the detection limit (<0.0040 mg/kg) to 22 mg/kg. The soil vapor analyses also showed similar measurements of BTEX and TPH, with concentrations of TPH ranging from 280 ppmv to 19,000 ppmv and from 0.11 ppmv (ethylbenzene) up to 67 ppmv (total xylenes) of the BTEX compounds (Table 2). The results of the soil chemistry analyses are summarized in Table 3.

## 2.2.2 Soil Gas Permeability and Radius of Influence

The raw data for the soil gas permeability test at Building 1639 are presented in Appendix C. Using the Hyperventilate<sup>TM</sup> computer model, soil gas permeabilities were calculated at each of the monitoring points. These data are presented in Table 4. The soil gas permeability varied considerably, with values ranging from 24 darcy up to 5.4 x 10<sup>8</sup> darcy. Typically, the radius of influence is calculated by plotting the log of the pressure change at a specific monitoring point versus the distance from the vent well. The radius of influence would then be the distance where 1 inch of water pressure can be measured. However, in this instance, 1 inch of water pressure was not achieved at any monitoring point (Figure 5); therefore, a radius of influence based on these specifications cannot be definitively determined at this site, other than to say it is less than 8.7 feet.

## 2.2.3 In Situ Respiration Test

The results of the in situ respiration test for Building 1639 are presented in Appendix D. Each figure in Appendix D illustrates the oxygen, carbon dioxide, and helium concentrations as a

Table 2. Results From Soil and Soil Gas Analyses for BTEX and TPH at Building 1639

Matrix	Sample Name	Benzene (mg/kg)	Toluene (mg/kg)	Ethylbenzene (mg/kg)	Total Xylenes (mg/kg)	TPH¹ (mg/kg)
Soil	H1-VW-4'-4.5'	0.048	0.020	0.015	0.020	22
	H1-VW-4.5'-5.0'	0.67	0.27	0.43	0.45	15
	H1-A-3'-4'	1.0	4.3	1.3	12	< 0.0040
Matrix	Sample Name	Benzene (ppmv)	Toluene (ppmv)	Ethylbenzene (ppmv)	Total Xylenes (ppmv)	TPH² (ppmv)
Soil Gas	H1-A-2.5	5.2	4.2	1.1	2.9	5,600
	H1-A-5	27	35	10	30	19,000
	H1-B-2.5	2.8	1.3	0.84	1.9	2,700
	H1-B-5	2.4	0.84	0.42	1.5	3,200
	H1-C-3.5	0.44	0.13	0.11	0.37	280
	H1-C-6	11	20	9.3	67	11,000

Referenced to a reference oil composed of a mixture of 2,2,4-trimethylpentane, *n*-hexadecane, and chlorobenzene.

<sup>&</sup>lt;sup>2</sup> TPH referenced to jet fuel (molecular weight = 156).

Table 3. Results From Soil Chemistry Analyses at Building 1639

	Sample Name				
Parameter	H1-VW-4'-4.5'	H1-VW-4.5'-5.0'	H1-A-3'-4'		
Alkalinity (mg/kg CaCO <sub>3</sub> )	< 50	< 50	< 50		
Moisture (% by weight)	22.2	21.8	5.8		
рН	5.8	6.0	6.1		
Iron (mg/kg)	7,980	6,260	8,630		
Total Phosphorous (mg/kg)	370	290	300		
Total Kjeldahl Nitrogen (mg/kg)	1,100	730	70		
Particle Size Analysis (%)	Gravel: 11.5	Gravel: 0.5	Gravel: 26		
	Sand: 59	Sand: 72	Sand: 55		
	Silt: 27	Silt: 23	Silt: 16		
	Clay: 6	Clay: 4.5	Clay: 3		

Table 4. Results of Hyperventilate™ Soil Gas Permeability Analysis at Building 1639

Monitoring Point	Depth (ft)	Soil Gas Permeability (darcy)
Н1-МРА	2.5	24
	5.0	130
Н1-МРВ	2.5	1,400
	5.0	6,200
Н1-МРС	3.5	5.4 x 10 <sup>8</sup>
	6.0	28

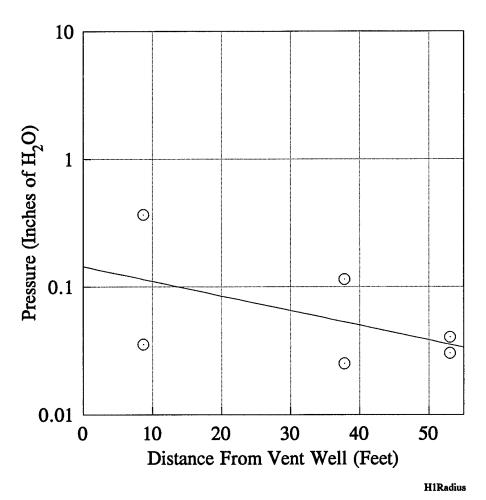


Figure 5. Radius of Influence at Building 1639

function of time. An example of typical oxygen utilization at this site is shown in Figure 6, where oxygen utilization and carbon dioxide production at monitoring point H1-MPA-5.0' is illustrated. A summary of the oxygen utilization and carbon dioxide production rates and corresponding biodegradation rates is shown in Table 5. The biodegradation rates measured at this site were fairly high, with rates ranging from 8.0 mg/kg/day to 27 mg/kg/day based on oxygen utilization, and from 0.48 mg/kg/day to 4.3 mg/kg/day based on carbon dioxide production.

Loss of helium was insignificant at all monitoring points, indicating that the monitoring points were well sealed and that the oxygen depletion observed was a result of biodegradation.

Soil temperatures were measured during the in situ respiration test. Temperatures during the test ranged from 17.9 to 19.3°C at monitoring point H1-MPA-2.5′ and from 18.7 to 20.7°C at monitoring point H1-MPA-5.0′.

## 2.2.4 Bioventing Demonstration

The decision was made to install a bioventing system at Building 1639. A 1-HP blower was installed at the site on October 14, 1992. Air injection was initiated on October 14 at a flowrate of 2.5 scfm.

Table 5. Oxygen Utilization and Carbon Dioxide Production Rates During the In Situ Respiration Test at Building 1639

Monitoring Point	Oxygen Utilization Rate (%/hour)	Biodegradation Rate (mg/kg/day)	Carbon Dioxide Production Rate (%/hour)	Biodegradation Rate (mg/kg/day)
Background	0.0073	0.14	0.0087	0.19
H1-MPA-5.0'	1.4	27	0.076	1.6
H1-MPB-2.5'	0.42	8.0	0.034	0.74
H1-MPB-5.0'	0.58	11	0.022	0.48
H1-MPC-6.0'	0.69	13	0.20	4.3

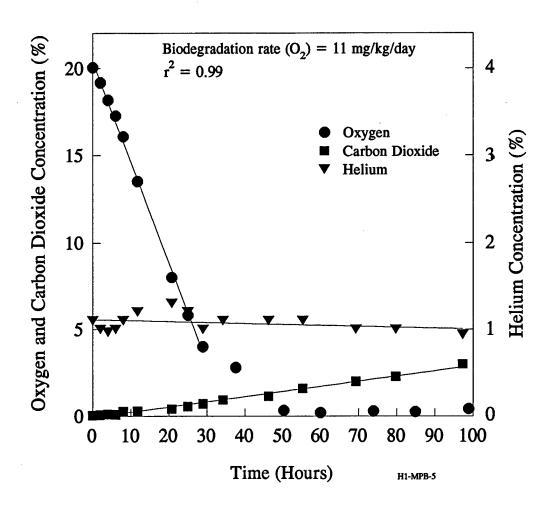


Figure 6. Oxygen Utilization and Carbon Dioxide Production During the In Situ Respiration Test at Monitoring Point H1-MPB-5.0'

## 3.0 BUILDING 1812

## 3.1 Chronology of Events and Site Activities

Although oxygen levels measured during the soil gas survey indicated that some areas at the Building 1812 site may be limited, oxygen concentrations measured at the permanent monitoring points were fairly high. Therefore, an in situ respiration test was not conducted at this site. However, because measurements taken during the soil gas survey indicated that some areas were oxygen-limited, a bioventing system was installed at the site in order to treat those areas. Other activities were conducted at the site according to the Test Plan and Technical Protocol (Hinchee et al., 1992).

#### 3.1.1 Groundwater Measurements

Groundwater depth was measured at one monitoring well (HB-02) at Building 1812. The groundwater level was measured on October 1, 1992 and was recorded at 3.98 feet.

## 3.1.2 Soil Gas Survey

On October 1, 1992, a limited soil gas survey was conducted to locate a suitable test area at Building 1812. Soil gases were sampled by driving a %-inch-diameter stainless steel probe into the soil with a hammer drill. Soil gas was withdrawn with a vacuum pump and analyzed for oxygen, carbon dioxide, and TPH. Soil gas measurements were taken as described in Section 2.1.2.

The soil gas probes were driven to depths ranging from 2.0 to 5.0 feet at several locations at Building 1812. Table 6 provides the initial concentrations of oxygen, carbon dioxide, and TPH for the various locations at Building 1812. Relatively high concentrations of oxygen were found at most of the soil gas probes, with concentrations ranging from 3 to 20%. Relatively low concentrations of carbon dioxide (0.05 to 8.5%) and TPH (10 ppm to 800 ppm) were encountered. The oxygen concentrations in the soil gas indicate that some areas at this site are oxygen-limited and may respond to bioventing.

Table 6. Initial Soil Gas Composition at Building 1812

Monitoring Point	Depth (ft)	Oxygen (%)	Carbon Dioxide (%)	TPH (ppm)
GS-1	2.5	19.5	0.7	120
GS-2	2.5	20	0.05	10
	5.0	NS	NS	NS
GS-3	2.25	20.2	0.60	87
GS-4	2.5	17	0.30	40
	4.0	19.8¹	0.08	20
GS-5	2.5	15	3.8	220
	5.0	13	5.1	270
GS-6	2.5	16.5	1.3	- 120
	5.0	NS	NS	NS
GS-7	2.0	9.1	6.0	440
	3.5 - 4.0	7.0	7.2	480
GS-8	2.5	17.5	1.2	250
GS-9	2.0	17	1.4	280
GS-10	2.0	3.0	8.5	800
GS-11	2.0	18.8	0.40	180
	3.0	18.3	0.60	96
	4.0	18.9¹	2.7	220
GS-12	2.0	17	0.10	41
	3.0	18	0.70	120
	4.0	17.5	0.10	50

NS Not sampled. Groundwater was encountered at this depth.

Pressure reading on sampling pump was high. Measured oxygen concentration may not be representative of actual soil gas oxygen concentrations. The actual oxygen concentration is likely to be lower.

## 3.1.3 Vent Well, Monitoring Point, and Thermocouple Installation

On October 4, 1992, one vent well and three monitoring points were installed, and soil samples were collected for analyses. The monitoring points were labeled H2-MPA, H2-MPB, and H2-MPC. The locations of the vent well and monitoring points are shown in Figure 3. A cross section of the vent well and monitoring points showing site lithology and construction detail is shown in Figure 7.

The vent well was installed at a depth of 7.0 feet into an 8-inch-diameter borehole. The vent well consisted of Schedule 40 2-inch-diameter PVC piping with 3.6 feet of ten-slot screen. The annular space corresponding to the screened area of the well was filled with silica sand, whereas the annular space above the screened interval was filled with bentonite to prevent short-circuiting of air to or from the surface.

Soil gas probes consisted of ¼-inch tubing with a 1-inch-diameter, 6-inch screened area. The annular space corresponding to the screened area was filled with silica sand, whereas the interval between the screened areas was filled with bentonite, as was the annular space from the shallowest monitoring point to the ground surface. The monitoring points were installed at depths as follows:

- Monitoring point H2-MPA was installed at a depth of 5.0 feet into an 8-inch-diameter borehole. The monitoring point was screened to two depths: 2.5 and 5.0 feet.
- Monitoring point H2-MPB was installed at a depth of 7.0 feet into an 8-inch-diameter borehole. The monitoring point was screened to three depths: 2.5, 5.0, and 7.0 feet.
- Monitoring point H2-MPC was installed at a depth of 6.0 feet into an 8-inch-diameter borehole. The monitoring point was screened to three depths: 2.5, 4.5, and 6.0 feet.

A Type K thermocouple was installed with monitoring points H2-MPA-2.5' and H2-MPA-5.0'.

MPC

MPB

MPA

Vent Well

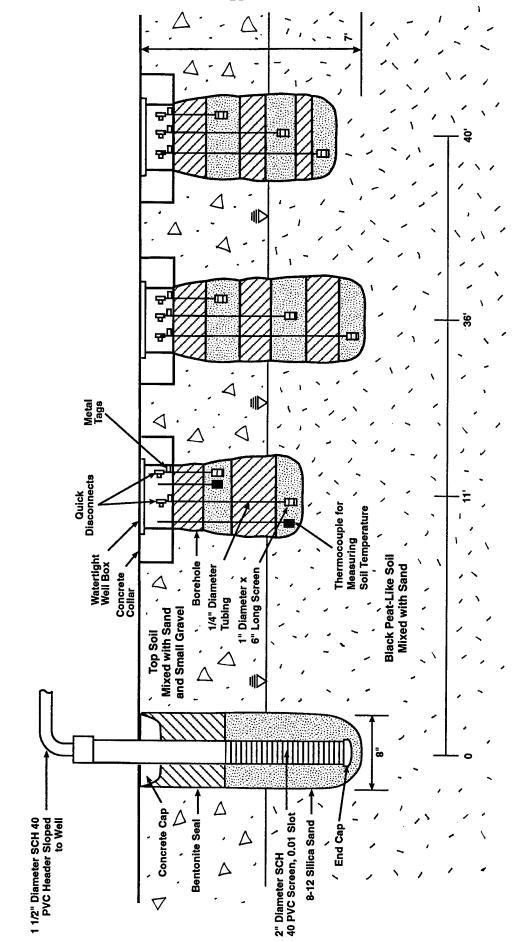


Figure 7. Cross Section of Vent Well and Monitoring Points at Building 1812 Showing Site Lithology and Construction Detail (not to scale)

F/Kittel11/h-2

## 3.1.4 Soil and Soil Gas Sampling and Analyses

Split-spoon soil samples were collected at depths of 3.0 to 3.5 feet and 4.0 to 4.5 feet from the vent well borehole and were labeled H2-VW-3'-3.5' and H2-VW-4'-4.5', respectively. The samples were sent under chain of custody to Engineering-Science Inc., Berkeley Laboratory for analyses of BTEX; TPH; alkalinity; moisture content; pH; iron; total phosphorous; total Kjeldahl nitrogen; and particle size analysis.

## 3.1.5 Soil Gas Permeability and Radius of Influence

A detailed description of the method for conducting a soil gas permeability test, including equations to compute k, the soil gas permeability, is given in the Test Plan and Technical Protocol (Hinchee et al., 1992).

Prior to air injection, the monitoring points were allowed to set up for 24 hours. A portable 1-HP explosion-proof positive displacement blower unit was used to inject air. After air injection was initiated, pressure readings were taken approximately every 1 to 2 minutes for the first hour, then approximately every 10 minutes for the following hour. The Hyperventilate<sup>TM</sup> computer model was used to calculate the soil gas permeability.

## 3.2 Results and Discussion

## 3.2.1 Soil and Soil Gas Analyses

Results of the soil analyses for BTEX and TPH are presented in Table 7. The analytical report for this site is presented in Appendix B. All of the BTEX compounds were at concentrations below the detection limit in sample H1-VW-3'-3.5' and only small quantities of toluene (0.015 mg/kg) and total xylenes (0.051 mg/kg) were detected in sample H1-VW-4'-4.5'. TPH concentrations were low in sample H1-VW-3'-3.5' (12 mg/kg); however, 13,000 mg/kg of TPH was detected in sample H1-VW-4'-4.5'. The results of the soil chemistry analyses are summarized in Table 8.

Table 7. Results From Soil Analyses for BTEX and TPH at Building 1812

Sample Name	Benzene (mg/kg)	Toluene (mg/kg)	Ethylbenzene (mg/kg)	Total Xylenes (mg/kg)	TPH¹ (mg/kg)
H2-VW-3'-3.5'	< 0.062	< 0.072	< 0.052	< 0.093	12
H2-VW-4'-4.5'	< 0.0032	0.015	< 0.0026	0.051	13,000

Referenced to a reference oil composed of a mixture of 2,2,4-trimethylpentane, *n*-hexadecane, and chlorobenzene.

Table 8. Results From Soil Chemistry Analysis at Building 1812

	Sample Name				
Parameter	H2-	-VW-3'-3.5'	H2-VW-4'-4.5'		
Alkalinity (mg/kg CaCO <sub>3</sub> )		90		< 50	
Moisture (% by weight)		3.2		6.1	
рН	7.0			6.2	
Iron (mg/kg)	8,160			5,460	
Total Phosphorous (mg/kg)		450		600	
Total Kjeldahl Nitrogen (mg/kg)		66		53	
Particle Size Analysis (%)	Gravel:	10	Gravel:	1.0	
	Sand:	69.5	Sand:	76	
	Silt:	19	Silt:	21.5	
	Clay:	1.5	Clay:	1.5	

## 3.2.2 Soil Gas Permeability and Radius of Influence

The raw data for the soil gas permeability test at Building 1812 are presented in Appendix E. Using the Hyperventilate<sup>TM</sup> computer model, soil gas permeabilities were calculated at each of the monitoring points. These data are presented in Table 9. The soil gas permeability varied considerably, with values ranging from 2.5 darcy up to 6.3 x 10° darcy. The radius of influence where 1 inch of water could be measured was calculated by plotting the log of the pressure change at the monitoring points versus the distance from the vent well (Figure 8). If pressure changes at all monitoring points are included, no radius of influence can be calculated based on these specifications. However, if the pressure change at monitoring points below the water table are not included (H2-MPA-5.0' and H2-MPB-5.0'), the radius of influence at Building 1812 is estimated to be approximately 7.5 feet.

## 3.2.3 Bioventing Demonstration

The decision was made to install a bioventing system at Building 1812. A 1-HP blower was installed on October 14, 1992. Air injection was initiated on October 14 at a flowrate of 4.25 scfm.

## 4.0 BACKGROUND AREA ACTIVITIES

The background area was located as shown in Figure 1. An existing monitoring well was used as the vent well and was screened from 1.5 feet to 9.0 feet. Soil samples were taken 2 feet from the monitoring well by hand auger. Site lithology at this area was representative of that in the contaminated areas.

A split-spoon soil sample was collected at a depth of 2.5 to 3.0 feet from the vent well borehole and was labeled H1-BKG-2.5'-3'. A soil vapor sample also was collected from the vent well after installation and labelled H1-BG-1.5-9. The soil samples were sent under chain of custody to Engineering-Science Inc., Berkeley Laboratory for analysis of BTEX; TPH; alkalinity; moisture content; pH; iron; total phosphorous; total Kjeldahl nitrogen; and particle size analysis. The soil vapor sample was sent under chain of custody to Air Toxics, Ltd., in Rancho Cordova, California, for analysis of BTEX and TPH.

Table 9. Results of Hyperventilate™ Soil Gas Permeability Analysis

Monitoring Point	Depth (ft)	Soil Gas Permeability (darcy)
H2-MPA	2.5	6.3 x 10°
	5.0	2.5
Н2-МРВ	2.5	3.1 x 10°
	5.0	430
	7.0	ND
H2-MPC	2.5	NR
	4.5	NR
	6.0	NR

ND No data were collected from this monitoring point.

NR No pressure readings were detected at this monitoring point.

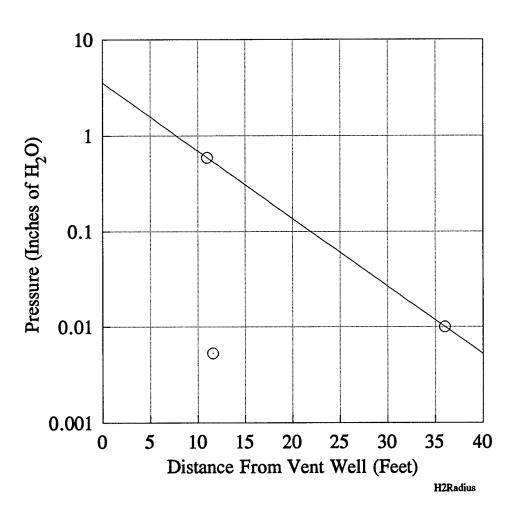


Figure 8. Radius of Influence at Building 1812

Results of the soil and soil vapor analyses for BTEX and TPH are presented in Table 10. The analytical report for the background area is presented in Appendix B. All of the BTEX compounds were at concentrations below the detection limit in the soil sample, and only a small quantity of TPH (84 mg/kg) was detected. The soil vapor sample contained higher concentrations of BTEX compounds with concentrations ranging from 0.16 ppmv (benzene) up to 4.7 ppmv (total xylenes). The results of the soil chemistry analyses also are summarized in Table 10.

An in situ respiration test was conducted at the background area beginning on October 9 after 24 hours of air injection. The test was concluded on October 11. Very little decrease in oxygen concentration occurred during the course of the in situ respiration test (Figure 9).

#### 5.0 FUTURE WORK

Base personnel will be required to perform a simple weekly system check to ensure that the blower is operating within its intended flowrate, pressure, and temperature range. An on-site briefing for base personnel who will be responsible for blower system checks was conducted when the blowers were installed. The principle of operation was explained, and a simple checklist and logbook were provided for blower data. Base personnel will be asked to perform minor maintenance activities, such as replacing filters or gauges, or draining condensate from knockout chambers, but they will not be expected to perform complicated repairs or analyze gas samples. Replacement filters and gauges will be provided and shipped to the base, and serious problems, such as motor or blower failures, will be corrected by Battelle.

The progress of this system will be monitored by conducting semiannual respiration tests in the vent well and in each monitoring point and by regularly measuring the oxygen, carbon dioxide, and hydrocarbon concentrations in the extracted soil gas and comparing them to background levels. At least twice each year, the progress of the bioventing test will be reported to the base point-of-contact.

Table 10. Results From Soil Chemistry Analysis at Background Area

	Sample Name	
Parameter	Soil Sample H1-BKG-2.5'-3'	Soil Vapor Sample H1-BG-1.5-9
Benzene	<0.00060 mg/kg	0.16 ppmv
Toluene	<0.00070 mg/kg	0.93 ppmv
Ethylbenzene	<0.00050 mg/kg	0.43 ppmv
Total Xylenes	<0.00090 mg/kg	4.7 ppmv
ТРН	84 mg/kg	340 ppmv
Alkalinity (mg/kg CaCO <sub>3</sub> )	<50	
Moisture (% by weight)	9.8	
pН	6.6	
Iron (mg/kg)	11,600	
Total Phosphorous (mg/kg)	460	
Total Kjeldahl Nitrogen (mg/kg)	91	
Particle Size Analysis (%)	Gravel: 11.5	
	Sand: 48	
	Silt: 32	
	Clay: 8.5	

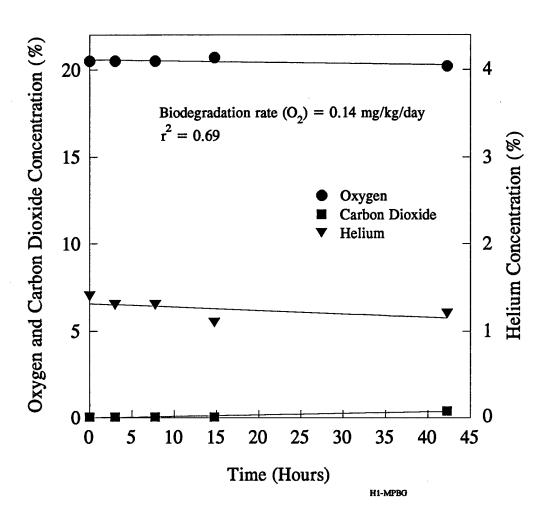


Figure 9. Oxygen Utilization and Carbon Dioxide Production During the In Situ Respiration Test at the Background Area

## 6.0 REFERENCE

Hinchee, R.E., S.K. Ong, R.N. Miller, D.C. Downey, and R. Frandt. 1992. *Test Plan and Technical Protocol for a Field Treatability Test for Bioventing* (Rev. 2), Report prepared by Battelle Columbus Operations, U.S. Air Force Center for Environmental Excellence, and Engineering Sciences, Inc. for the U.S. Air Force Center for Environmental Excellence, Brooks Air Force Base, Texas.

## APPENDIX A

TEST PLAN FOR HANSCOM AFB, MASSACHUSETTS



505 King Avenue Columbus, Ohio 43201-2693 Telephone (614) 424-6424 Facsimile (614) 424-5263

September 2, 1992

Captain Catherine Vogel
Department of the Air Force
Building 1117
HQ AFESC/RDVW
Tyndall AFB, Florida 32403-6001

Dear Cathy:

SUBJECT: TEST PLAN FOR BIOVENTING INITIATIVE FIELD TEST AT BUILDING 1639 AND BUILDING 1812, HANSCOM AFB, MA.

This letter was prepared to accompany the report titled "Test Plan and Technical Protocol for a Field Treatability Test for Bioventing." The report was developed as a generic test plan for the Air Force Bioventing Initiative Project in which Hansom AFB is participating. This letter outlines site specific information to support the generic test plan.

The sites chosen for the bioventing test initiative are Building 1639 and Building 1812. Building 1639 is the base fuel service station (see map Figure 1). There have been three reported petroleum releases (gasoline, #2 fuel oil, and waste lubricating oil) at this facility since November 1990. The site at Building 1812 is the location of a heating oil fuel tank (see Figure 2). The tank was replaced recently and soil TPH concentrations as high as 2830 mg/Kg have been reported.

The purpose of this project is to investigate the feasibility of using the bioventing technology to remediate petroleum contaminated soils at the above mentioned sites.

#### **Site Description:**

Hanscom AFB is located in Bedford, Massachusetts. A generalized geologic and hydrogeologic column is shown in Figure 3. Groundwater at Hanscom is encountered at 3-8 feet. Figures 4 and 5 show test boring logs for two soil borings in the vicinity of the bioventing initiative sites.

Soil contamination at Building 1639 ranges from 599 ppm to 4400 ppm TPH. Soil samples at Building 1812 have indicated TPH concentrations as high as 2800 ppm. Boring and depth locations for these samples were not immediately available. Locations for initiating the soil gas survey at each site will be determined with direction from the Base POC.

Captain Catherine Vogel
Department of the Air Force
September 2, 1992
Page 2

#### Project Activities-

The following field activities are planned for the bioventing project at Hanscom AFB. The same procedures will be followed at each site. Additional detail can be found in Section 5.0 of the test plan and technical protocol.

- A small scale soil gas survey will be conducted to identify an appropriate location for installation of the bioventing system. The soil gas survey will be conducted in areas which site data have shown to be the most contaminated. Soil vapor from the candidate site should exhibit high petroleum hydrocarbon concentrations (10,000 ppm or greater), relatively low O<sub>2</sub> concentrations (0 % to 2.0 %), and relatively high CO<sub>2</sub> concentrations (depending on soil type, 2.0 % to 10.0 %, or higher). An uncontaminated background location will also be identified.
- Once the installation sites are located one vent well and three 3-level soil gas monitoring points will be installed in the contaminated location and one vent well will be installed in a background area (one background area will be used for both test sites, if possible). The wells and monitoring points will be installed using a two-man power auger or a portable drill rig to bore down to just above the water table. Three to four soil samples will be collected for chemical/physical analysis.
- 3- The air permeability test will be conducted in the contaminated test location.
- 4- Following the air permeability test, in situ respiration tests will be conducted in both the contaminated and the background test locations.
- Depending on the results of the air permeability test and the in situ respiration test, a decision will be made whether or not to install a blower system in the contaminated area for the long term bioventing test. If the decision is made to install the system, the blower will be plumbed to the vent well and bioventing will be started (assuming power is available). Site personnel will be trained for blower operation prior to Battelle leaving the site.
- A report detailing the results of the in situ respiration test and the air permeability test will be provided to the project officer and the base POC.

#### Schedule-

Field activities at Hanscom AFB are planned to begin on September 28, 1992. Battelle will have 2 to 3 people on site for approximately 3 weeks.

Captain Catherine Vogel Department of the Air Force September 2, 1992 Page 3

#### Base Support-

Hanscom AFB needs to be able to provide the following:

- Digging permits and utility clearance need to be obtained prior to the initiation of the field work. Underground utilities should be clearly marked to reduce the chance of utility damage or personal injury during soil gas probe and well installation. Battelle will not be able to begin field operations without these clearances.
- Electrical power will need to be easily accessible from the project site. The air permeability test and in situ respiration test can be performed using a gasoline powered electric generator. The operation of the bioventing system will require a permanent 220/110 V power source. If power will not be available immediately after the test is completed the bioventing system will be installed for start-up at a later date.
- The Air Force will need to provide drums to contain soil cuttings and provide for contaminated soil disposal.
- Base and site clearance will be required for Battelle's site employees. We will furnish you with personal information for each person at least one week prior to starting field operations.
  - Regulatory approval, if any is required, will need to be obtained by the base prior to start-up of the bioventing system. The system will likely be configured for air injection so there will be no point source vapor emission from the system. The wells to be installed will not intersect the apparent water table and no groundwater will be pumped. A letter from the Massachusetts DEP (Attachment 1) presented several concerns with the bioventing test plan. These concerns are addressed below.

#### Regulatory Issues-

- General Approval Requirements- DEP approval is necessary for the initiation of remediation of petroleum releases. This approval should be pursued by the base for the long-term bioventing test. The soil gas survey, vent well and monitoring point installation, air permeability test, and in situ respiration test are all standard field activities that generally do not require any regulatory approval prior to initiation. Battelle is available to discuss any of these activities with the DEP prior to start up of field activities.
- Air Quality Permits and Approvals- The only air injection planned for this study is the injection of ambient atmospheric air into the vadose zone as a oxygen source for biological activity. Should conditions require the extraction of soil gas (i.e. a basement adjacent to the vent well) then treatment and permitting requirements will be discussed with the DEP.

Captain Catherine Vogel Department of the Air Force September 2, 1992 Page 4

- Groundwater Remediation- The purpose of this research project is to investigate the efficacy of bioventing for the remediation of petroleum contaminated soils in the vadose zone. While bioventing, as configured for this project, does not directly address groundwater contamination, it has proven to be an effective technology for the remediation of petroleum contaminated soils. These soils typically act as the long-term source of localized groundwater contamination. Removal of the contaminant source is paramount in any groundwater remediation effort.
- 4- Nutrient Addition- Nutrient addition is not currently planned for this site.
- 5- Field Screening of Split-Spoon Samples- The Jar Headspace Technique will be employed for selected samples. Samples will be analyzed for BTEX via EPA Method 8020 and TPH via EPA Method 418.1. Samples will also be analyzed for nutrients.

Please let me know if there are any other regulatory concerns which need to be addressed. If you have any questions please feel free to call me at (614) 424-6122.

Sincerely,

Jeffrey A. Kittel
Researcher
Environmental Technology

JAK:mla

# ATTACHMENT 1



Commonwealth of Massachusetts

Executive Office of Environmental Affairs

# Department of Environmental Protection

Metro Boston/Northeast Regional Office:

William F. Weld Governor Daniel S. Greenbaum Commissioner

JUN 1 0 1992

Mr. Robert Spelfogel Hanscom Air Force Base 3245th ABG/DEEV Bedford, MA 01730

RE: BEDFORD - Hanscom AFB
Bioventing Treatability Test
DEP Case #3-3882

Dear Mr. Spelfogel:

The Department of Environmental Protection is in receipt of a January 1992 "Report, Test Plan And Technical Protocol For A Field Treatability Test For Bioventing". This report was prepared for the U.S. Air Force by Battelle Corporation and was submitted to the Department by the Air Force with a request to provide information concerning the regulatory requirements of the bioventing remediation technique. Preliminary approval of the bioventing concept was also requested. Hanscom Air Force Base is planning to contract Battelle Corporation to conduct Bioventing Treatability Testing at Buildings 1812 and 1639.

Building 1639 (DEP Case #3-3882) is the Air Force Base Service Station. Department files indicate three petroleum release incidents at this location. These releases were assigned the incident response numbers N92-0837, N91-794 and N90-1884. Gasoline, #2 fuel oil, and waste lubricating oil have been released to soil and/or groundwater at this site.

At Building 1812 soil samples collected from borings were found to be contaminated with concentrations of Total Petroleum Hydrocarbons as high as 2830 mg/kg.

The bioventing process uses aeration of subsurface soils to stimulate in-situ biological activity and promote bioremediation. The following points outline the bioventing process.

# Hanscom AFB Page 2

- (1) Forced air is supplied to the contaminated unsaturated zone to produce the aerobic conditions necessary for biodegradation.
- (2) Airflow is optimized to reduce volatilization while maintaining aerobic conditions.
- (3) Local soil gas conditions are monitored to assure aerobic conditions.
- (4) Moisture and nutrients are added to the contaminated unsaturated zone if necessary.
- (5) Dewatering is conducted, if necessary to lower the water table below the contaminated soils.

After review of the above mentioned report the Department hereby approves of the initiation of Bioventing Field Treatability Testing at Buildings 1639 and 1812. The following comments are also provided:

\* General Approval Requirements.

Departmental approval must be obtained for the initiation of the remediation of releases of petroleum or hazardous materials at Locations To Be Investigated or Confirmed Disposal Sites. A bioventing system to remediate petroleum or hazardous materials releases at Hanscom Air Force Base is subject to the above-mentioned approvals.

Air Quality Permits and Approvals.

Page 5 Section 2.1.3 states that when air is injected into a contaminated zone and withdrawn from clean soils volatile hydrocarbons are allowed to degrade prior to being withdrawn thereby eliminating contaminated off-gases. "This configuration typically does not require air emission permitting".

Be advised that all air emissions resulting from treatment systems at petroleum or hazardous materials disposal sites are subject to the Air Pollution Control Regulations at 310 CMR 6.00-8.00 in addition to the requirements of the Massachusetts Contingency Plan at 310 CMR 40.000.

Hanscom AFB Page 3

In general, if an air contaminant source emits less than one ton of contaminants per year, then that emission source is not subject to application for approval from the Department of Environmental Protection's Division of Air Quality.

Whether or not the emission source emits more than one ton of pollutants per year, if the emission is part of a system being operated to remediate a release of petroleum or hazardous materials to the environment, then that system is subject to M.G.L. c.21E and the MCP. The DEP Bureau Of Waste Site Cleanup requires that off-gas controls be installed on all point source remedial air emissions where Soil Vapor Extraction systems are utilized. Off-gas controls must be implemented during the first 180 days of operation at a minimum. Off-gas treatment must reduce vapor-phase emissions of VOC's by at least 95 percent. Approval to operate without emission controls after the first 180 days are contingent upon receipt of information confirming the absence of a threat of harm to human health or the environment.

Page 5 indicates that during a 1988 study at Hill Air Force Base it became apparent that bioventing had great potential for remediating JP-4 fuel-contaminated soils.

It is not clear however that bioventing is effective for remediation of groundwater contaminated with fuel oils. Many of the components contained in aircraft fuels, diesel fuel and heavier petroleum fuel products are not significantly volatile and would not readily be transferred from the groundwater to the unsaturated zone where they could be treated by the bioventing system. In order to achieve a permanent solution to the petroleum contamination present at a site it may be necessary to reduce the levels of contaminants in soil and groundwater to levels which do not pose a significant or otherwise unacceptable risk to public health, safety, public welfare or the environment.

Page 6 indicates that adding nutrients as required to increase biodegradation rates may be necessary. The addition of nutrients to the subsurface is subject to the requirements of M.G.L. c.21 and the Groundwater Discharge Permit Regulations (314 CMR 5.00). Nutrient addition may be initiated only upon receipt of a Groundwater Discharge Permit issued by the DEP Division of Groundwater Pollution Control.

Hanscom AFB Page 4

Page 42, Section 2.1.2 - Exploratory Boring in Deep Soils, states that "Split-spoon samples will be visually checked for fuel contamination and screened for volatile emissions by passing a hydrocarbon analyzer slowly over the open split-spoon".

This method is not acceptable for the field screening of split spoon samples for petroleum hydrocarbon contamination. The Jar Headspace Technique (Reference: "Management Procedures for Excavated Soils Contaminated with Virgin Petroleum Oils", Policy #WSC-89-001) should be applied for the measurement of volatile petroleum constituents. Total Petroleum Hydrocarbons by Infrared (Standard Methods 503 or 5520) or Oil Fingerprinting (ASTM D 3328) should be used to determine concentrations of non-volatile petroleum hydrocarbons.

If you have any questions regarding this matter please do not hesitate to contact Jack Miano at the letterhead address or telephone (617) 935-2160 X142.

Very truly yours,

Jack Miano

Environmental Engineer

Stephen M. Johnson

Acting Chief,

Site Management Branch

SMJ/JM/ae

cc: DEP, BWSC, Boston, Attn: Jeff Krukonis

DEP, DWS, NERO, Attn: Jim Persky

Bedford BOH, Attn: David Black

Bedford DPW, 312 Great Rd., Bedford, MA 01730,

Attn: Mr. Richard Warrington, Director

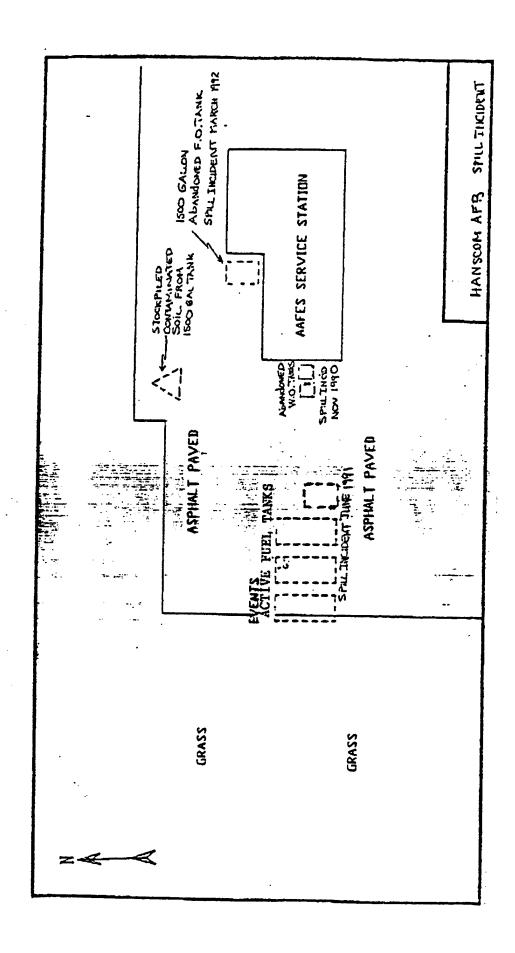


FIGURE 1. SITE DIAGRAM FOR BUILDING 1639

# FORMER LOCATION OF UNDERGROUND STORAGE TANK BUILDING NO. 1816

HANSCOM AIR FORCE BASE, MASSACHUSETTS

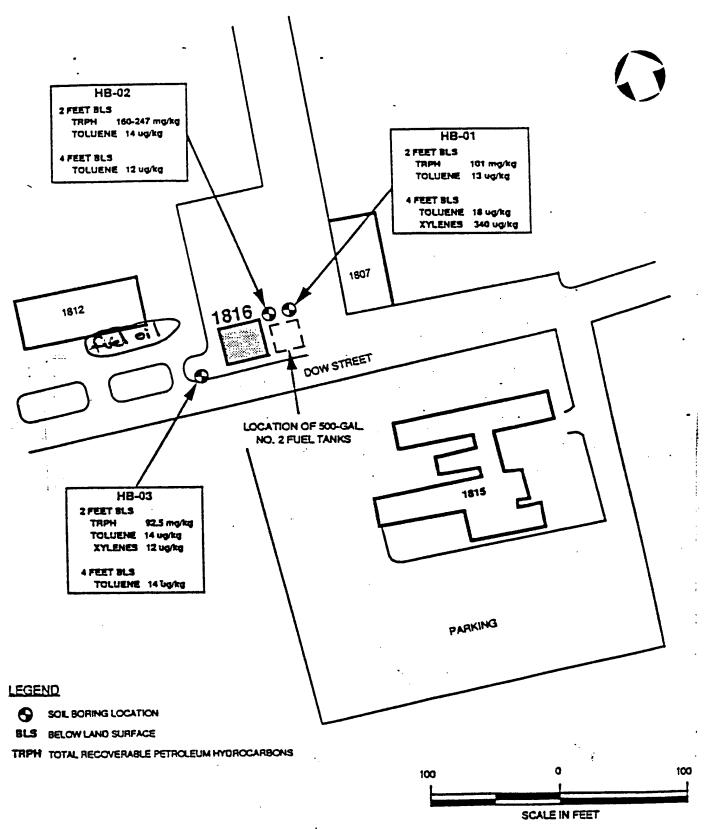


FIGURE 2. SITE DIAGRAM FOR BUILDING 1812

# GENERALIZED GEOLOGIC AND HYDROGEOLOGIC COLUMN INVESTIGATION OF SUSPECTED HAZARDOUS WASTE SITES HANSCOM AIR FORCE BASE, MASSACHUSETTS

PERIOD	ЕРОСН	FORMATION	COLUMNAR SECTION	APPROXIMATE THICKNESS	CHIARACTER		
	RECENT	FILL —unconformable—		0-10	TOPSOIL & CLEAN SANDY FILL		
	1	PEAT —unconformable—		0-7	BLACK ORGANIC SANDS & PEAT		
QUATERNARY	(NIS	? OUTWASH DEPOSITS		0-18	Clean Coarse to fine Sands SATURATED, PERMEABLE	UPPER AQUIFER	
	PLEISTOCENE (WISCONSIN)	PLEISTOCENE (WISCO	PLEISTOCENE (WISCO	LACUSTRINE DEPOSITS		0-65	Fine SANDS & SILTS Grading to dayey Silts, Low permeability, SATURATED
		Sub-Lacustrine Deposits —unconformable—	مم	0-20	Fine Sands & Silts, Saturated Permeable		
		GLACIAL TILL —unconformable—		0-10	Coarse to fine SAND, trace to some SILT, trace to some GRAVEL	LOWER AQUIFER	
		ANDOVER GRANITE		0-?	Medium to Coarse Grained Muscovite- Biotite Granite	BEDROCK AQUIFER	

BORING NUMBER	HB-19	
JOB NUMBER	11-9540	
DATE STARTED	4-24-90	
DATE COMPLETED	4-24-90	
DRILLED BY	TN	
LOGGED BY	KMP	
CHECKED BY	GPM	

#### REMARKS:

PAGE 1 OF 1

FLUSH MOUNTED WELL
SWL MEASURED 9.34' BELOW TOC ON 5/29/90
WELL ALIGNMENT TEST COMPLETED 5/18/90
HNU READINGS RECORDED IN "LAB TEST" COLUMN

ELEV. IN FEET	DEPTH IN FEET	DESCRIPTION	MONITORING WELL CONSTRUCTION	SYM- BOLS	LAB TESTS	SPT N VALUE
129.02	0.0	Firm yellowish brown (10 YR 5/6) to very pale brown (10 YR 7/3) fine to medium grained SAND (SP)			0	19
-		·		·	0	22 -
122.02		Firm light brownish gray (2.5 YR 6/2) fine grained SAND (SP)			. 0	17
-		Very stiff light yellowish brown (2.5 Y 6/4) SILT (ML)		<u>_</u>	0	16 -
117.02	12.0	Very stiff light yellowish brown (2.5 Y 6/4) SILT with lenses of fine grained sand (ML)			<b>0</b>	17
113.52	15.5	Boring Terminated at 15.5				16 -
_		·			·	
-						-
		•,				
-					•	-

FIGURE 4. TEST BORING RECORD FOR SOIL BORING IN THE VICINITY OF BUILDING 1812

BORING NUMBER	HB-03	
JOB NUMBER	11-9540	
DATE STARTED	5-1-90	
DATE COMPLETED	5-1-90	
DRILLED BY	TN	
LOGGED BY	KMP	
CHECKED BY	GPM	

REMARKS:

PAGE 1 OF 1

FLUSH MOUNTED WELL SWL MEASURED 3.05 BELOW TOC ON 5/29/90 WELL ALIGNMENT TEST COMPLETED 5/18/90 HNU READINGS RECORDED IN "LAB TEST" COLUMN

1		Ţ				
	ELEV. IN FEET	DEPTH IN FEET	DESCRIPTION	MONITORING WELL SYM- CONSTRUCTION BOLS	LAB TESTS	SPT N VALUE
,	132.00	0.0	Very stiff to hard light olive brown (2.5 Y 5/4) SILT (ML)	<b>Y</b>	0	32
					0	20 —
	123.00	9.0			0	<b>27</b>
	120.00	12.0	Firm yellowish brown (10 YR 5/6) fine to medium grained SAND with stratifications (SM)		0	17 -
		13.0	Very stiff grayish brown (2.5 Y 5/2) clayey SILT (ML)  Boring Terminated at 12.0°		0	31
						-
	-	:				
			.*			
	-	·				_
		·				
H	-					4
L						1

# APPENDIX B

ANALYTICAL REPORT FOR BUILDING 1639, BUILDING 1812, AND BACKGROUND AREA

# (a) AIR TOXICS LTD.

AN ENVIRONMENTAL ANALYTICAL LABORATORY

# WORK ORDER #: 9210075

Work Order Summary

CLIENT:

Mr. Jeff Kittel

BILL TO:

Accounts Payable

Battelle

Engineering Science

505 King Ave.

1700 Broadway Ste. 900

Columbus, OH 43201

Denver, CO 80290

PHONE:

614-424-6122

**INVOICE # 8630** 

FAX:

614-424-3667

**P.O.** # DE268.03.04

DATE RECEIVED:

10/14/92

**AMOUNT:** \$910.00

DATE REPORTED:

10/26/92

**PROJECT #** G4468-0640

			Receipt	
FRACTION #	<u>NAME</u>	<u>TEST</u>	VAC./Press.	PRICE
01A	H1-A-2.5	TO-3	0 "Hg	\$120.00
02A	H1-A-5	TO-3	0 "Hg	\$120.00
03A	H1-B-2.5	TO-3	2.0 "Hg	\$120.00
04A	H1-B-5	TO-3	1.0 "Hg	\$120.00
05A	H1-C-3.5	TO-3	0.5 "Hg	\$120.00
06A	H1-C-6	TO-3	0.5 "Hg	\$120.00
07A	H1-BG-1.5-9	TO-3	1.0 "Hg	\$120.00
0 <b>7</b> B	H1-BG-1.5-9 Duplicate	TO-3	1.0 "Hg	NC
08A	Method Spike	TO-3	NA	NC
09A	Lab Blank	TO-3	NA	NC
Misc Charges	1 Liter SUMMA Canister P	reparation (7)	@ \$10.00 each.	<b>\$7</b> 0.00

CERTIFIED BY Amold & Frances

Laboratory Director

DATE: 10/96/93

SAMPLE NAME: H1-A-2.5 ID#: 9210075-01A

## **EPA Method TO-3**

(Aromatic Volatile Organics in Air)

## BTXE BY GC/PID

File Name: Dil. Factor:	610150		Date of Collection	
Dit. Pactor.	MDL	MDL	Date of Analysis: Amount	10/15/92 Amount
Compound	(ppmv)	(uG/L)	(ppmv)	(uG/L)
Benzene	0.20	0.62	5.2	16
Toluene	0.20	0.74	4.2	15
Total Xylenes	0.20	0.85	2.9	12
Ethyl Benzene	0.20	0.85	1.1	4.7

# TOTAL PETROLEUM HYDROCARBONS GC/FID

File Name; Dil. Factor;	610150 <b>2</b> 0		Date of Collection  Date of Analysis:	
Compound	MDL (ppmv)	MDL (uG/L)	Amount (ppmv)	Amount (uG/L)
TPH*	2.0	8.0	5600	22000

<sup>\*</sup>TPH referenced to Jet Fuel (MW=156)

SAMPLE NAME: H1-A-5 ID#: 9210075-02A

## **EPA Method TO-3**

(Aromatic Volatile Organics in Air)

## BTXE BY GC/PID

File Name: Dil. Factor:	610150 50		Date of Collection Date of Analysis:	
	MDL	MDL	Amount	Amount
Compound	(ppmv)	(uG/L)	(ppmv)	(uG/L)
Benzene	0.50	1.6	27	84
Toluene	0.50	1.8	35	130
Total Xylenes	0.50	2.1	30	130
Ethyl Benzene	0.50	2.1	10	42

# TOTAL PETROLEUM HYDROCARBONS GC/FID

File Name: 6101509 Date of Collection: 10/12/92 Dil. Factor: 500 Date of Analysis: 10/15/92					
	MDL	MDL	Amount	Amount	
Compound	(ppmv)	(uG/L)	(ppmv)	(uG/L)	
TPH*	5.0	20	19000	76000	

<sup>\*</sup>TPH referenced to Jet Fuel (MW=156)

SAMPLE NAME: H1-B-2.5 ID#: 9210075-03A

## **EPA Method TO-3**

(Aromatic Volatile Organics in Air)

# BTXE BY GC/PID

File Name: 6101510 Date of Collection: 10/12/92						
Dil. Factor:	22	0	Date of Analysis:	10/15/92		
	MDL	MDL	Amount	Amount		
Compound	(ppmv)	(uG/L)	(ppmv)	(uG/L)		
Benzene	0.22	0.69	2.8	8.7		
Toluene	0.22	0.81	1.3	4.8		
Total Xylenes	0.22	0.93	1.9	8.1		
Ethyl Benzene	0.22	0.93	0.84	3.6		

# TOTAL PETROLEUM HYDROCARBONS GC/FID

File Name:	610151		Date of Collection	
Dil. Factor:	22	0	Date of Analysis:	10/15/92
	MDL	MDL	Amount	Amount
Compound	(ppmv)	(uG/L)	(ppmv)	(uG/L)
TPH*	2.2	8.8	2700	11000

<sup>\*</sup>TPH referenced to Jet Fuel (MW=156)

SAMPLE NAME: H1-B-5 ID#: 9210075-04A

## **EPA Method TO-3**

(Aromatic Volatile Organics in Air)

# **BTXE BY GC/PID**

File Name:	610160	16	Date of Collection	1: 10/12/92
Dil. Factor:	21	0	Date of Analysis:	10/16/92
	MDL	MDL	Amount	Amount
Compound	(ppmv)	(uG/L)	(ppmv)	(uG/L)
Benzene	0.21	0.66	2.4	7.5
Toluene	0.21	0.77	0.84	3.1
Total Xylenes	0.21	0.89	1.5	6.4
Ethyl Benzene	0.21	0.89	0.42	1.8

# TOTAL PETROLEUM HYDROCARBONS GC/FID

File Name:	610160	6	Date of Collection	ı: 10/12/92
Dil. Factor:	21	0	Date of Analysis:	10/16/92
	MDL	MDL	Amount	Amount
Compound	(ppmv)	(uG/L)	(ppmv)	(uG/L)
TPH*	2.1	8.4	3200	13000

<sup>\*</sup>TPH referenced to Jet Fuel (MW=156)

SAMPLE NAME: H1-C-3.5 ID#: 9210075-05A

## **EPA Method TO-3**

(Aromatic Volatile Organics in Air)

# BTXE BY GC/PID

File Name:	610160	7	Date of Collection	n: 10/12/92
Dil. Factor:	1	0	Date of Analysis:	10/16/92
	MDL	MDL	Amount	Amount
Compound	(ppmv)	(uG/L)	(ppmv)	(uG/L)
Benzene	0.010	0.031	0.44	1.4
Toluene	0.010	0.037	0.13	0.48
Total Xylenes	0.010	0.042	0.37	1.6
Ethyl Benzene	0.010	0.042	0.11	0.47

# TOTAL PETROLEUM HYDROCARBONS GC/FID

File Name:	610160	17	Date of Collection	n: 10/12/92
Dil. Factor:	1	0	Date of Analysis:	10/16/92
	MDL	MDL	Amount	Amount
Compound	(ppmv)	(uG/L)	(ppmv)	(uG/L)
TPH*	0.10	0.40	280	1100

<sup>\*</sup>TPH referenced to Jet Fuel (MW=156)

SAMPLE NAME: H1-C-6 ID#: 9210075-06A

## **EPA Method TO-3**

(Aromatic Volatile Organics in Air)

## BTXE BY GC/PID

File Name: 6101514 Date of Collection: 10/12/92 Dil. Factor: 2100 Date of Analysis: 10/15/92				
	MDL	MDL	Amount	Amount
Compound	(ppmv)	(uG/L)	(ppmv)	(uG/L)
Benzene	2.1	6.6	11	34
Toluene	2.1	7.7	20	74
Total Xylenes	2.1	8.9	67	<b>2</b> 80
Ethyl Benzene	2.1	8.9	9.3	39

# TOTAL PETROLEUM HYDROCARBONS GC/FID

File Name:	610151	4	Date of Collection	n: 10/12/92
Dil. Factor:	210	0	Date of Analysis:	10/15/92
	MDL	MDL	Amount	Amount
Compound	(ppmv)	(uG/L)	(ppmv)	(uG/L)
TPH*	21	84	11000	44000

<sup>\*</sup>TPH referenced to Jet Fuel (MW=156)

SAMPLE NAME: H1-BG-1.5-9 ID#: 9210075-07A

## **EPA Method TO-3**

(Aromatic Volatile Organics in Air)

# BTXE BY GC/PID

File Name: Dil. Factor:	610160 1	14 0	Date of Collection Date of Analysis:	
	MDL	MDL	Amount	Amount
Compound	(ppmv)	(uG/L)	(ppmv)	(uG/L)
Benzene	0.010	0.031	0.16	0.50
Toluene	0.010	0.037	0.93	3.4
Total Xylenes	0.010	0.042	4.7	20
Ethyl Benzene	0.010	0.042	0.43	1.8

# TOTAL PETROLEUM HYDROCARBONS GC/FID

File Name; Dil. Factor:	610160	4 0	Date of Collection	
Compound	MDL (ppmv)	MDL (uG/L)	Date of Analysis: Amount (ppmv)	Amount (uG/L)
TPH*	0.10	0.40	330	1300

<sup>\*</sup>TPH referenced to Jet Fuel (MW=156)

## 9210075 Battelle

# AIR TOXICS LTD.

SAMPLE NAME: H1-BG-1.5-9 Duplicate ID#: 9210075-07B

## **EPA Method TO-3**

(Aromatic Volatile Organics in Air)

# BTXE BY GC/PID

File Name:	6101605 Date of Collection: 10/12/92				
Dil. Factor:	1	0	Date of Analysis:	10/16/92	
	MDL	MDL	Amount	Amount	
Compound	(ppmv)	(uG/L)	(ppmv)	(uG/L)	
Benzene	0.010	0.031	0.16	0.50	
Toluene	0.010	0.037	0.93	3.4	
Total Xylenes	0.010	0.042	4.7	20	
Ethyl Benzene	0.010	0.042	0.43	1.8	

# TOTAL PETROLEUM HYDROCARBONS GC/FID

File Name: Dil. Factor:	610160 1		Date of Collection  Date of Analysis:	
Compound	MDL (ppmv)	MDL (uG/L)	Amount	Amount
TPH*	0.10	0.40	( <b>ppmv)</b> 340	(uG/L) 1400
	0.10	0.10	040	1400

<sup>\*</sup>TPH referenced to Jet Fuel (MW=156)

SAMPLE NAME: Method Spike ID#: 9210075-08A

## **EPA Method TO-3**

(Aromatic Volatile Organics in Air)

## BTXE BY GC/PID

File Name: Dil. Factor:	610160 1.		Date of Collection: 10/12/92 Date of Analysis: 10/16/92
	MDL	MDL	***************************************
Compound	(ppmv)	(uG/L)	% Recovery
Benzene	0.001	0.003	90
Toluene	0.001	0.004	83
Total Xylenes	0.001	0.004	80
Ethyl Benzene	0.001	0.004	80

# TOTAL PETROLEUM HYDROCARBONS GC/FID

File Name: Dil. Factor:	610160 1.		Date of Collection: 10/12/92 Date of Analysis: 10/16/92
	MDL	MDL	
Compound	(ppmv)	(uG/L)	% Recovery
TPH*	0.010	0.040	110

<sup>\*</sup>TPH referenced to Jet Fuel (MW=156)

SAMPLE NAME: Lab Blank ID#: 9210075-09A

# **EPA Method TO-3**

(Aromatic Volatile Organics in Air)

# BTXE BY GC/PID

File Name: Dil. Factor:	610150 1.		Date of Collection Date of Analysis:	
***************************************	MDL	MDL	Amount	Amount
Compound	(ppmv)	(uG/L)	(ppmv)	(uG/L)
Benzene	0.001	0.003	Not Detected	Not Detected
Toluene	0.001	0.004	Not Detected	Not Detected
Total Xylenes	0.001	0.004	Not Detected	Not Detected
Ethyl Benzene	0.001	0.004	Not Detected	Not Detected

# TOTAL PETROLEUM HYDROCARBONS GC/FID

File Name: Dil. Factor:	610150 1.		Date of Collection Date of Analysis:	
	MDL	MDL	Amount	Amount
Compound	(ppmv)	(uG/L)	(ppmv)	(uG/L)
TPH*	0.010	0.040	Not Detected	Not Detected

<sup>\*</sup>TPH referenced to Jet Fuel (MW=156)



11325 SUNRISE GOLD CIRCLE, SUITE 'E' RANCHO CORDOVA, CA 95742 (916) 638-9892 • FAX (916) 638-9917

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# CHAIN OF CUSTODY RECORD

•	MAT/TA		
	COLLECTED BY (Signature	ייין אין אין אין אין אין אין אין אין אין	
	TO0		
: (	#O.#		
34468 - 01.40			
) # TOBI DBG	BEMADKS # -	ריים פעניעושון: ו	

VAC /PRESSIBE	LAD I.D. #		0,4%	2.0"#J	1"14%	051145		**************************************	1.44	<b>3</b>	
ANALYSIS	DITE! 1-11	DICK IVH	-	=		(1)	=			<i>j</i>	
DATE/TIME	120ct 10900	200	=   -				-		+*************************************		
FIELD SAMPLE I.D.# SAMPLING MEDIA (Tenax, Canister etc.)	H-H-C.J (ANISEL	5.4.1x	HI-8-2.5 II	1-0-0	41-6-3.5	11-1-1		11 82-15-8			

DATE/TIME RECEIVED BY: DATE/TIME				CONDITION	
IED BY: DATE/TIME RELINQUISHED BY: DATE/TIME	08/2	LAB USE ONLY	OPENED BY DATERIAL		
RECEIVANISHED BY: DATE/TIME RECEIV	HODAM		HIPPER NAME AIR BILL #		SEMARKS

BERKELEY LABORATORY 600 BANCROFT WAY BERKELEY, CA 94710 Tel: (415) 841-7353

Report Date: November 19, 1992

Work Order No.:4432

Client:

Doug Downey

ES Denver/AFCEE/Hanscom AFB

1700 Broadway Denver, CO 80290

Date of Sample Receipt: 10/05/92

Your soil samples identified as:

H1-VW-4'-4.5' H1-VW-4.5'-5.0' H1-A-3'-4' H2-VW-3'-3.5' H2-VW-4'-4.5' H1-BKG-2.5'-3'

were analyzed for BTEX by EPA Method 8020, TRPH by EPA Method 418.1, TKN, total phosphorus, soil classification, pH, alkalinity, iron and moisture.

The analytical reports for the samples listed above are attached.

GC VOLATILES DATA PACKAGE

# VOLATILE ORGANICS CASE NARRATIVE WORK ORDER NO. 4432 EPA METHOD 8020

Sample H1-VW-S $^3$ -3.5' (4432-04) was run as a medium level due to the presence of high concentrations of non-target analytes.

## GC AMALYTICAL REPORT Mnalytical Detino BTEX Aromatic Compounds

Work Order NO.:4452

% Moisture: 22.21

Client ID:HI-UW-4/-4.5/

Matrix: SUIL

Laboratory ID:4452-1

Level:LOW

Date Collected: 10/02/92

Unit:UG/KG

Dilution Factor:

Date Analyzed:10/15/92 Date Confirmed:NA

Compound	Result	Reporting Limit	
			THULE
Senzene	48.0 D=5	0.8	
Ethyl Benzene	17.0	Ŭ. o	t <i>•</i>
fulyene	20.0	0.9	
Xylenes (total)	20.0	1.2	

ND-Not Detected NH-Not Applicable D-Dilution Factor

HHALYST: AS

GROUP LEADER: hur

## GC ANALYTICAL REPORT Analytical Method BIEX Aromatic Compounds

Work Order NO.:4432

% Moisture: 21.81

Client ID:HI-UW-4.5/-5.0/

Matrix:SUIL

Laboratory ID:4432-2

Level:MEDIUM

Date Collected: 10/02/92

Unit:UG/KG

Dilution Factor:

Date Analyzed:10/14/92

Date Confirmed:NA

 Compound	Result	Reporting Limit
Benzene	670.0	77.0
Ethyl Benzene	450.0	64.0.
Tolhene	270.0	90.0
Xylenes (total)	450.0	120.0

ND-Not Detected NA-Not Applicable D-Dilution Factor

ANHLYST: AG

GROUP LEADER : Lugar

# GC ANALYTICAL REPORT Analytical Method BTEX Aromatic Compounds

Work Order NO.:4432

Client ID:HI-A-3'-4'

Laboratory ID:4432-3

Date Collected: 10/02/92

Dilution Factor:

% Moisture: 5.81

Matrix:SOIL

Level:MEDIUM

.. . ....

Unit:UG/KG

Date Analyzed:10/14/92

Date Confirmed:NA

Compound	Result	Reporting Limit	
Benzene	1000.0	130.0	
Ethyl Benzene	1300.0	110.0	
Totuene	4500.0	150.0	
Xylenes (total)	12000.0	190.0	

ND-Not Detected NA-Not Applicable D-Dilution Factor

ANALYST: #B

GROUP LEADER : how

## GC ANALYTICAL REPORT Hnalutical Method BTEX Aromatic Compounds

Work Order NO.:4452 Order NO.:4432 Client ID:HZ-VW-3'-3.5'

% Moisture: 3.2

Matrix: SUIL

Laboratory 1D:4432-4

Level:MEDIUM

Date Collected: 10/03/92

Unit:UG/KG

Dilution Factor:

Date Analyzed:10/14/92 Date Confirmed:NA

Compound	Result	Reporting Limit
Benzene	NO	52.0
Ethyl Benzene	· ND	52.0
Toluene	ND	72.0
Xylenes (total)	ND	93.0

ND-Not Detected NA-Not Applicable D-Dilution Factor

ANALYST: AS

GROUP LEADER: Kusse

## GC ANALYTICAL REPORT Analytical Method BTEX Aromatic Compounds

Work Order NO. : 4452 TP 11/18/92

Client ID:H2-UW-4/-4.5/

Laboratory ID:4452-5

Date Collected: 10/05/92

Dilution Factor:

F-.

% Moisture: 6.14

Matrix:50IL

Level:LOW

Unit:UG/KG

Date Analyzed:10/15/92

Date Confirmed:NA

Compound	Result	Reporting Limit
Benzene	ND	3 . Z
Ethyl Benzene	ทบ	2.6
ïuluene	15.0	3.7
Xylenes (total)	51.0	4.8

ND-Not Detected NA-Not Applicable D-Dilution Factor

ANALYST: AS

GROUP LEADER: Rever

#### GC ANALYTICAL REPORT Hnalutical Method BIEX Aromatic Compounds

Work Order NO.:4432

Client ID:HI-BkG-2.5/-5/

% Moisture:NA

Matrix:SOIL

Laboratory ID:4432-7

Level:LOW

Date Collected: 10/04/92

Unit:UG/KG

Date Analyzed:10/13/92

Dilution Factor:

Date Confirmed:NA

Compound	Result	Reporting Limit	
			====
Benzene	ND	9.6	
Ethyl Benzene	Nū	0.5	
totuene	ND	0.7	
Xvlenes (total)	NÜ	0.9	

ND-Not Detected NA-Not Applicable D-Dilution Factor

ANALYST: AD

# GC ANALYTICAL REPORT Analytical Method BIEX Aromatic Compounds

Work Order NO.:4432

% Moisture:NA

Client ID:METHOD BLANK

Matrix:SOIL

Laboratory ID:MSVG4921015

Level:LOW

Date Collected: NA

Unit:UG/KG

Dilution Factor:

Date Analyzed:10/15/92 Date Confirmed:NA

Compound Result Reporting Limit Benzene NO 0.6 Ethyl Benzene ND 0.5 Toluene ΝÜ. 0.7 Xylenes (total) ND 0.9

ND-Not Detected NA-Not Applicable D-Dilution Factor

ANALYST: AD

GROUP LEADER : fund

### GC ANALYTICAL REPORT Hnalutical Method BIEX Aromatic Compounds

Work Order NO.:4452

% Moisture:NA

Client ID:METHOD BLANK

Matrix: SOIL

Laboratory ID:MWUG4921014

Level:MEDIUM

Date Collected: NA

Unit:UG/KG

Dilution Factor: 1

Date Analyzed:10/14/92

Date Confirmed:NA 

Compound	Result	Reporting Limit	<b> </b>
Benzene	ND	60.0	
Ethyl Benzene	ND	50.0	
toluene	NU	70.0	
Xylenes (total)	ND	90.0	

ND-Not Detected NA-Not Applicable D-Dilution Factor

ANALYST: AS

GROUP LEADER: Limi

# GC AMALYTICAL REPORT Headytical Dethod BTEX Aromatic Compounds

Work Order NO. #4452

Client ID: METHOD BLANK

Matrix:50IL

·

1001 IX-0011

Laboratory ID:M5VG4921015

Level:LOW

Date Collected: NA

Unit:UG/KG

Dilution Factor:

Date Analyzed:10/15/92

% Moisture:NA

Date Confirmed:NA

m: m;	Compound	Kesult	Reporting Limit			
	Benzene	ND	0.6			
	Ethyl Benzene	ND	Ü.5			
	Toluene	NŪ	0.7			
	Xylenes (total)	ND	0.9			

ND-Not Detected NA-Not Applicable D-Dilution Factor

ANALYST: #5

GROUP LEADER: L

ES-ENGINEERING SCIENCE, INC.

LABORATORY NO.

600 BANCROFT WAY BERKELEY, CA 94710

# GC ANALYTICAL REPORT ANALYTICAL REPORT BTEX AROMATIC COMPOUNDS

MATRIX: SOIL

DATE: 10/14/92

a-a-a-TrıFluoro

• • • • •		
MWUG4921014	METHOD BLANK	. yl
4452-2	HI-UW-4.57-5.07	0.2
4432-5	H1-A-3 -4'	115
4452-4	H-2-UW-5/-3.5	122

CLIENT ID

600 BANCROFT WAY BERKELEY, CA 94710

# GC ANALYTICAL REPORT -ANALYTICAL REPORT BTEX AROMATIC COMPOUNDS

MATRIX: SOIL

DATE:10/13&15/92

LABORATORY NO.	CLIENT ID	a-a-a-TriFluoro Toluene
MSUG4921013	METHOD BLANK	94
4432-7	HI-BKG-2.5'-3'	114
4435-1	KAFB66-SB1-SS3-5-5.5	108
MSVG4921015	METHOD BLANK	79
4432-1	HI-UW-4/-4.5/	118
4452-1DIL	HI-VW-41-4.51DIL	104
4432-5	H-F-VW-41-4.51	7.0
4451-1	HZ-UW-41-4.51 MC-UW-7	109
	TP 11/18	

# OUALITY CONTROL RESULTS SUMMARY ANALYTICAL REPORT BTEX AROMATIC COMPOUNDS

Work Order No.: 4432,4451,4455

QC sample No.: 55VG4921009A&B

Date analuzed:10/09/92

Matrix: SOIL

Dilution factor:1

			=====:		=====	====			
	•	SR UG/KG	i hs		I   MSD  UG/KG	•	RED	IOC L: I IRPD	IMITS       PR
COMPOUND     8020 analysis	   S#    UG/KG	SR UG∕KG	l MS	•	   MSD  UG/KG	•	I  RPD	QC   L :     RPD	[MIT5       PR
********	======	======			=====		=====		
Benzene     Toluene	20       20		22.6    21.4	İ	i	į	İ	į	39-150   
!	· · · · · · · · · · · · · · · · · · ·		1	ļ.	ł	l	1	[	1
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MS = Spike sample

MSD = Spike sample duplicate

SR = Sample result

SH = Spike added

ND = Not Found At or Above Detection Limits

 $RPD = 100 \times (MS-MSD)/((MS+MSD)/2)$ 

 $PR = 100 \times ((MS \text{ or } MSD) - SR)/SA$ 

ANALYST: AT

ac: M

= Not calculated

= Not Applicable

= Out of limits

NC

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# QUALITY CONTROL RESULTS SUMMARY ANALYTICAL REPORT BTEX AROMATIC COMPOUNDS

Work Order No.: 4432,4451,4435

QC sample No.: SWVG4921016A&B Date analyzed:10/16/92

Matrix: SOIL Dilution factor:1

**************************************		SR UG/KG	MS	•	I   MSD  UG/KG		=====    RPD 	QC L     RPD	IMITS
COMPOUND     8020 analysis	   SA    UG/KG	SR UG∕KG	MS	IMS IPR	I I MSD IUG/KG		  RPD 	IQC L:     RPD	IMITS
Benzene     Toluene 	İ			1	Ì	82   82    135 	20     4 	į į	39-150   

MS = Spike sample

MSD = Spike sample duplicate

SR = Sample result

SA = Spike added

ND = Not Found At or Above Detection Limits

 $RPD = 100 \times (MS-MSD)/((MS+MSD)/2)$ 

 $PR = 100 \times ((MS \text{ or } MSD) - SR)/SA$ 

ANALYST: A

ac: pws

= Not calculated

= Not Applicable

= Out of limits

NC

NA

### WO # 4432,4451,4435

LAB NAME : ENGINEERING-SCIENCE, INC. DATE ANALYZED 10/13&15/92

LAB SAMPLE ID:MSUG4921013&15

DATE EXTRACTED : NA

MATRIX :LOW SOIL

INSTRUMENT ID: VGC-4

LAB	CLIENT	DATE
SAMPLE ID	SAMPLE ID	ANALYZED
MSUG4921013	METHOD BLANK	10/13/92
4432-7	HI-BKG-2.5/-3/	10/13/92
4435-1	KAFB66-SB1-SS3-5-5.51	10/13/92
MSVG492101 🔩	METHOD BLANK	10/15/92
4432-1	HI-UW-4'-4.5'	10/15/92
4432-1DIL	HI-VW-4'-4.5'DIL	10/15/92
4432-5	HE-UW-41-4.51	10/15/92
4451-1	MC-VW-7	10/15/92
t e e e e e e e e e e e e e e e e e e e	TP 11/18	
	117	

### METHOD BLANK SUMMARY

WU # 4432

LAB NAME : ENGINEERING-SCIENCE, INC. DATE ANALYZED :10/14/92

LAB SAMPLE ID:MWVG4921014

DATE EXTRACTED : NA

MATRIX :SOIL

INSTRUMENT ID: VGC-4

LAB SAMPLE ID	CLIENT Sample ID	DATE ANALYZED
MWUG4921014 4432-2 4432-3 4432-4	METHOD BLANK HI-VW-4.5/-5.0/ HI-A-3/-4/ Ht-VW-3/-3.5/	10/14/92 10/14/92 10/14/92 10/14/92
	40 11/18	

TOTAL RECOVERABLE PETROLEUM HYDROCARBONS

DATA PACKAGE

### ORGANIC ANALYTICAL REPORT

Work Order NO.: 4432

Parameter: TPH

Matrix: Soil

Unit: mg/Kg

Analytical

Method: 418.1

Date Extracted 10/27/92

QC Batch NO.: S92QCB026TPH

Date Analyzed: 10/29/92

		化苯酚 的复数有效 医自然性 医性性性			
Sample ID:		Client ID:	Result	Reporting Limit	Percent Moisture
		<b>计算证明证据的证据的证据的现在记录的</b>			
4432-01		HI-VW-4'-4.5'	22	5	22.2
4432-02		HI-VW-4.5'-5.0'	<del>-</del> -		<del></del>
	,		15	5	21.8
4432-03		HI-A-3'-4'	ND	4	5.8
4432-04	18	HZ-VW-3'-3.5'	12	4	- <del>-</del>
4432-05	11/14			4	3.2
4432-05	141.	H2-VW-4'-4.5'	13000	4	6.1
4432-07		HI-BKG-2.5'-3'	84	5	11.4
MSTPH921027		— · <del>-</del> —		3	11.4
MS1FM32102/		METHOD BLANK	ИD	4	NA

NA\_ Not Analyzed ND\_ Not Detected

ANALYST:

Man D

GROUP, LEADER:

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# **INORGANICS DATA PACKAGE**

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6.2

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#### INORGANICS ANALYTICAL REPORT

Client: ES-Denver Work Order: 4432 Project: Matrix: Solid H₽→VW Client's ID: HI-BKG -3'-3.5' -2.5'-3' Sample Date: 10/03/92 10/03/92 10/04/92 % Moisture: Lab ID: 4432.04 4432.05 4432.06 Normal Parameter -----Results-----Method Report Units Date Limit Analyzed Alkalinity 90. ND ND SM 403(M) 50 mg/Kg CaCO3 11/02/92 3.2 Moisture 6.1 3.8 ASTM D2216 . 1 % by wt 10/22/92

6.6

EPA 9045

NA

pH Units

10/27/92

Note: Samples for alkalinity analysis were extracted using 10mL water for each 1g sample. These water extracts were analyzed for alkalinity, and the results were calculated in the solid on a dry-weight basis.

NA- Not Applicable ND- Not Detected

ANALYST: Don Deator

GROUP LEADER:

5.8

6. Đ

Moisture

'nď

10/22/92

10/27/92

. 1

NA

pH Units

#### INORGANICS ANALYTICAL REPORT

Work Order: Client: ES-Denver 4432 Matrix: Project: AFCEE Solid Client's ID: HI-VW HI-VW HI-A -4'-4.5' -4.5'-5.0' -3'-4' 10/02/92 Sample Date: 10/02/92 10/02/92 % Moisture: Lab ID: 4432.01 4432.02 4432.03 Normal Parameter -----Results-----Method Report Units Date: Limit Analyzed 50 ND ND ND SM 403(M) mg/Kg CaCO3 11/02/92 Alkalinity 5.8 22.2 21.8 ASTM D2216 \* by wt

6.1

EPA 9045

Note: Samples for alkalinity analysis were extracted using 10mL water for each 1g sample. These water extracts were analyzed for alkalinity, and the results were calculated in the solid on a dry-weight basis.

NA- Not Applicable ND- Not Detected

ANALYST:

GROUP LEADER:

600 Bancroft Way Berkeley, CA 94710

#### INORGANIC QC SUMMARY - MS and MSD

Work Order:

4432

% Moisture:

N A

Lab ID Spk/Dup:

Alkalinity Moisture pH

Matrix:

Solid

QC Batch:

Blank Spk 4438.04 4454.01 452.42 451.82 453.40

Units: mg/Kg CaCO3 (Alk)

\$ by wt. (Hois) pH Units (pH)

	Date Analyzed	Unspiked	Results		RPD -Conc Added- QC			Percent Recovered		
Parameter	MS/Dup	Sample	MS/Sample	MSD/Dup		Limit	NS	MSD	HS	MSD
Alkalinity Hoisture PH	11/02/92 10/22/92 10/27/92	0.00	23000.00 8.38 5.95	23000.00 7.52 6.08	0 11 2	20 29 26	23650.00	23650.00	97	וֿצ

\* or N = Outside QC Limit:

QC Limits for % Rec: 75 - 125

ANALYST: Don Steaton Date 11/11/92 REVIEWER: MWB Date 11/17/92. File: M1QCHSWH

#### INORGANICS ANALYTICAL REPORT

Client:

ES-Denver

Work Order:

4432

Project:

AFCEE

Matrix:

Solid

Client's ID:

Prep Blank

Sample Date:

% Moisture:

Lab ID:

Prep Blank

Parameter	Results	Method	Normal Report Limit	Units	Date Analyzed
Alkalinity	ND	SM 403(M)	50	mg/Kg CaCO3	11/02/92
Moisture	NA	ASTM D2216	.1	% by wt	10/22/92
рH	NA	EPA 9045	NA	pH Units	10/27/92

Note: Samples for alkalinity analysis were extracted using 10mL water for each 1g sample. These water extracts were analyzed for alkalinity, and the results were calculated in the solid on a dry-weight basis.

NA- Not Applicable ND- Not Detected

ANALYST:

Von Deston

GROUP LEADER:

### INORGANICS QC SUMMARY - LAB CONTROL SAMPLE

Work Order:

4432

% Moisture:

NA

Lab ID of LCS:

Alkalinity:

452.42 LCS

Matrix:

Solid

Units:

mg/Kg CaCO3

	Date Analyzed	LCS	Conc	% Rec	Advisory % Re	
Parameter	LCS	Result	Added	LCS	Low	High
Alkalinity	11/02/92	23000.00	23650.00	97	80	120

**METALS DATA PACKAGE** 

### CASE NARRATIVE WORK ORDER NO. 4432 METALS

The serial dilution sample result for iron did not agree with the undiluted result within 10%, and the diluted sample result was greater than ten times the iron MDL. All iron results in this batch are therefore flagged with "E".

Client ID's were abridged by the laboratory to facilitate computer entry of analytical data. The following should be used as a reference:

CLIENT ID		ABRIDGED ID
H1-VW-4'-4.5'		H1VW-4
H1-VW-4.5'-5.0'		VW-4.5
H1-A-3'-4'		A-3-4
H2-VW-3'-3.5'		VW-3
H2-VW-4'-4.5'	** * * · · · · · · · · · · · · · · · ·	H2VW-4
H1-BKG-2.5'-3'		BKG2.5

INORGANIO	ANALYSES DATA SHEET
Y_LABORATORY	H1VW-4   Contract: AFCEE_ES-D
Case No.: 4	32S SAS No.: SDG No.: A-3-4_
oir_	Lab Sample ID: 4432.01
OW	Date Sampled : 10/02/92
77.8	
ion Units (v	'L or mg/kg dry weight): MG/KG
Analyte	Concentration C Q M
-6 Iron	7980 E P
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FORM I - IN

CLIENT SAMPLE ID

•			,		CLIENT S	AMPLE ID
		INORGANIC .	ANALYSES DATA	SHEET		
Lab Name: E_S	BERKELEY_L	ABORATORY_	Contract: A	FCEE_ES-I	   VW-4	. 5
ab Code: ESB	L Ca	se No.: 44	32S SAS No.	:	SDG No.:	A-3-4_
atrix (soil/	water): SOIL	'_		Lab Samp	ole ID: 443	2.02
} Level (low/med	d): LOW_	_		Date Sam	mpled : 10/	02/92
Solids:	_78.	2				
Co	oncentration	Units (ug	/L or mg/kg dr	y weight)	): MG/KG	
	CAS No.	   Analyte	  Concentration	c	 	
			6260	1_1	_ii	
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•		INORGANIC	ANALYSES DATA S	SHEET		CL	JENT SAMPLE ID
Joh Nome E C	ז עם זפערפת		Contract. N		ת_ח		A-3-4
			Contract: Al			l	
Lab Code: ESBL	Ca	se No.: 44	32S SAS No.	<b>.</b>		SD	G No.: A-3-4_
Matrix (soil/w	ater): SOIL	<del></del>		Lab	Sampl	e I	D: 4432.03
Level (low/med	): LOW_	<del>_</del>		Date	Samp	led	1 : 10/02/92
% Solids:	_94.	2					
Co.	ncentration	Units (ug	/L or mg/kg dry	y wei	ght):	MG	:/KG
	CAS No.	Analyte	Concentration	C	Q	м	
·	7439-89-6	  Iron	8630	_     _     _	E!	P_	
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		INORGANIC	ANALYSES DATA	SHEET	СБТ	ENT SAMPLI	E 1D
						VW-3	1
ab Name: E_S_	BERKELEY_L	ABORATORY_	Contract: Al	FCEE_ES-	D i		i
ab Code: ESBI	Ca	se No.: 44	32S SAS No.	:	SDG	No.: A-3	-4_
atrix (soil/w	vater): SOIL	_		Lab Sam	ple ID	: 4432.04	
evel (low/med	l): LOW_			Date Sa	mpled	: 10/03/9	2
Solids:	_96.	8					
Co	oncentration	Units (ug	/L or mg/kg dr	y weight	): MG/	KG	
	CAS No.	   Analyte	  Concentration	ici ō	м		
	7439-89-6	Iron	8160	_ _E	P_		
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# INORGANIC ANALYSES DATA SHEET

CLIENT S.	AMPLE TO	)
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ab Name: E_S	BERKELEY_I	ABORATORY_	_ Contract: A	FCE	E_ES-	D	H2VW-	4
			32S SAS No.				G No.:	A-3-4_
latrix (soil/							D: 4432	
evel (low/med	d): LOW_						: 10/0	
Solids:	_93.	9						
Co	oncentration	Units (ug	//L or mg/kg dr	y w	eight	): MG	/KG	
-	CAS No.	   Analyte	Concentration	c	Q	М		
• •	7439-89-6	Iron	5460		E	_   _   _   P _	,	
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		INORGANIC	ANALYSES DATA	SHEET	CLIENT SAMPLE ID
Lab Name: E_S_	BERKELEY_L	ABORATORY_	Contract: A	FCEE_ES-D	BKG2.5
Lab Code: ESBL	Ca	se No.: 44	32S SAS No.	:	SDG No.: A-3-4_
Matrix (soil/w	ater): SOIL	<del>_</del>		Lab Samp	le ID: 4432.06
Level (low/med	): LOW_			Date Sam	pled : 10/04/92
Solids:	_90.	2			
Co	ncentration	Units (ug	/L or mg/kg dry	y weight)	: MG/KG
	CAS No.	   Analyte	Concentration	I C Q	М
	,	:	11600	E	   P_
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omments:					

		INORGANIC	ANALYSES DATA :	SHEET	CLI	ENT SAMPL	E ID
Lab Name: E S	BERKELEY L	ABORATORY	_ Contract: Al	FCEE ES-	     D	PBLANK	1
			32S SAS No.	_		No.: A-3	-4
Matrix (soil/w						: PBK 482	_
Level (low/med						: 11/04/9	
% Solids:	100.				•		_
Co	ncentration	Units (ug	/L or mg/kg dry	y weight	): MG/	KG	
,	CAS No.	   Analyte	  Concentration	I I ICI Q	   M		
	7439-89-6	l		l_l	_   _ P_		
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Comments:							_

## LABORATORY CONTROL SAMPLE (BLANK SPIKE)

lab Na	ame: E_S	BERKELEY_LABORATORY_	Contract:	AFCEE_ES	-D		
Lab Co	de: ESBL_	Case No.: 4432S	SAS No.:		SDG	No.:	A-3-4_
Solid	LCS Sourc	e: ESBL-LCSS					~
Aqueou	ıs LCS Sou	rce:					

Analyte	Aqueous (ug/L)   True Found %R			   True	its	878			
Iron			ļ	100.0	92.1		80.0	120.0	_92.1
		!							
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### LABORATORY CONTROL SAMPLE (BLANK SPIKE)

ab	Name:	E_SBERKI	ELEY_LABORATORY_	Contract:	AFCEE_ES	-D		
Lab	Code:	ESBL	Case No.: 4432S	SAS No.:		SDG	No.:	A-3-4_
ol	id LCS	Source:	ESBL-LCSS					
) qu	eous LO	CS Source:						

Analyte		eous (ug/I Found	!     True 	Solid Found C	l (mg/kg) Lim	%R	
Iron			 100.0	93.9 _	80.0	120.0	_93.9
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BLANK SPIKE DUPLICATE

CLIENT	SA	ME	L	E	ΙI	)
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			LCSSD
Lab Name: E_SBERKEL	EY_LABORATORY_	Contract: AFCEE_ES-D	•
Lab Code: ESBL	Case No.: 4432S	SAS No.:	SDG No.: A-3-4_
fatrix (soil/water):	soil_	Level	(low/med): _LOW
% Solids for Sample:	100.0	% Solids for	Duplicate: 100.0

Concentration Units (ug/L or mg/kg as received):MG/KG

    Analyte	Control     Limit		C	•	C	     RPD	ĺΩ	     M
Iron		92.0990	-	93.8830		1.9_	_   _	F_
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### ICP SERIAL DILUTION

EPA SAMPLE NO.

			1		
			1	H1VW-4L	
ontract:	AFCEE_	_ES-D	1		

Lab Name: E\_S\_BERKELEY\_LABORATORY\_ C

SDG No.: A-3-4\_

ab Code: ESBL\_\_\_ Case No.: 4432S\_ SAS No.: \_\_\_\_ atrix (soil/water): SOIL\_

Level (low/med): LOW\_\_\_

## Concentration Units: ug/L

	11	Serial	11 % 1	11
	Initial Sample	Dilution	Differ-	i i
Analyte	Result (I) C	Result (S)	C   ence	101
Iron	66454.92_ _	73844.92	_	EP
			_	- -
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## Engineering Science - Berkeley Laboratory

Method Detection Limits (Annually)

Lab Name:	E_SBERKE	LEY_LABORA	TORY_	Contract:	AFCEE_ES	5-D	
ab Code:	ESBL	Case No.:	4432S_	SAS No.:	•	S	SDG No.: A-3-4_
ICP ID Num	ber:	TJA_61_	м	Date:	08/31/92	2	
rlame AA I	D Number :			Matrix: S	sorr_		
urnace AA	ID Number	:		(ug/L in	1.00g to	100n	nl digestate)
) 	     Analyte	   Wave-     length     (nm)			MDL (ug/L)	М	·
	Iron	 271.44			46.0	   P	
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Comments:							

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### PREPARATION LOG

Lab Name: E\_S\_BERKELEY\_LABORATORY\_ Contract: AFCEE\_ES-D

Lab Code: ESBL\_\_\_ Case No.: \_4432S\_ SAS No.: \_\_\_\_ SDG No.:A-3-4\_

ethod: P\_

EPA   Sample   No.	  Preparation    Date	Weight   (gram)	Volume   (mL)
A-3-4	11/04/92	1.21	100
BKG2.5	11/04/92	1.23	100
1111111-4	11/04/92	1.07	
H2VW-4	1 11/04/92	1.24	100
LCSS	11/04/92		
LCSSD	11/04/92		
MPA-9	11/04/92		
MPB-9	11/04/92		
IPBLANK	11/04/92	1.00	100i
IVMP1-9	11/04/92		
VMP2-9	11/04/92		
VW-10	11/04/92		
VW-3	11/04/92	1.03	100
VW-4.5	- · <del></del>	1.18	100
VW - 7			
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#### ANALYSIS RUN LOG

\_Lab Name: E\_S\_\_BERKELEY\_LABORATORY\_ Contract: AFCEE\_ES-D

Lab Code: ESBL\_\_ Case No.: 4432S\_ SAS No.: \_\_\_\_ SDG No.:A-3-4\_

Instrument ID Number: TJA 61 M\_

Method: P\_

Start Date: 11/09/92

End Date: 11/09/92

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EPA	. [		l		!																						<del></del>	—
Sample	D/F	Time	ક	R	F			_ !		. !	- !	ļ	- [	. !	. !	- !	ļ	. !	!	[	. !	. !	- [		- [	Į	- !	
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### ANALYSIS RUN LOG

Lab Name: E\_S\_BERKELEY\_LABORATORY\_ Contract: AFCEE\_ES-D

Lab Code: ESBL\_\_\_ Case No.: 4432S\_ SAS No.: \_\_\_\_ SDG No.:A-3-4\_

Instrument ID Number: TJA 61 M\_

Method: P\_

Start Date: 11/09/92

End Date: 11/09/92

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ILMO2.1

TOTAL PHOSPHORUS

TOTAL KJELDAHL NITROGEN

SOIL CLASSIFICATION

DATA PACKAGE



Engineering Science, Inc. 600 Bancroft Way Berkeley, CA 94710

Attention: Tom Paulson

Client Project ID: Sample Descript:

Analysis for:

First Sample #:

W.O. #4432

Soil

Percent Solids 210-0637 Sampled:

9/2-4/92

Received: Analyzed:

Oct 7, 1992 Oct 19, 1992

Reported:

Oct 20, 1992

### LABORATORY ANALYSIS FOR:

### **Percent Solids**

	Sample Number	Sample Description	Detection Limit %	Sample Result %
	210-0637	H1-VW-4'-4.5'	10	83
	210-0638	H1-VW-4.5'-5.0'	10	84
,	210-0639	H1-A-S'-4'	10	94
	210-0640	H2-VW-3'-3.5'	10	97
	210-0641	H2-VW-4'-4.5'	10	92
	210-0642	H1-BKG-2.5'-3'	10	92

Analytes reported as N.D. were not present above the stated limit of detection.

**SEQUOIA ANALYTICAL** 

Tod Granicher Project Manager THIS REPORT HAS BEEN

APPROVED AND REVIEWED BY

ESBL PROJECT MANAGER

DATE

600 Bancroft Way

Berkeley, CA 94710 Attention: Tom Paulson Client Project ID: Sample Descript:

Analysis for:

First Sample #:

Soil

W.O. #4432

**Total Phosphorous** 

210-0637

Sampled:

9/2-4/92

Received: Analyzed:

Oct 7, 1992 Oct 19, 1992

Reported:

Oct 20, 1992

## LABORATORY ANALYSIS FOR:

# **Total Phosphorous**

Sample Number	Sample Description	Detection Limit mg/kg	Sample Result mg/kg	
210-0637	H1-VW-4'-4.5'	10	370	
210-0638	H1-VW-4.5'-5.0'	10	290	
210-0639	H1-A-3'-4'	10	300	
210-0640	H2-VW-3'-3.5'	10	450	
210-0641	H2-VW-4'-4.5'	10	600	
210-0642	H1-BKG-2.5'-3'	10	460	
• •	Method Blank	1.0	N.D.	

Analytes reported as N.D. were not present above the stated limit of detection.

**SEQUOIA ANALYTICAL** 

Tod Granicher **Project Manager**  Please Note:

Analysis results reported on a dry-weight basis.

2100637.ENG <8>

Engineering Science, Inc. 600 Bancroft Way

Berkeley, CA 94710 Attention: Tom Paulson Client Project ID: Sample Descript: W.O. #4432

Soil Total Kjeldahl Nitrogen

Analysis for: Total Kjeld First Sample #: 210-0637 Sampled:

9/2-4/92

Received: Analyzed: Oct 7, 1992 Oct 15, 1992

Reported: Oct 20, 1992

## LABORATORY ANALYSIS FOR:

# Total Kjeldahl Nitrogen

Sample Number	Sample Description	Detection Limit mg/kg	Sample Result mg/kg
210-0637	H1-VW-4'-4.5'	20	1,100
210-0638	H1-VW-4.5'-5.0'	20	730
210-0639	H1-A-3'-4'	20	70
210-0640	H2-VW-3'-3.5'	20	66
210-0641	H2-VW-4'-4.5'	20	53
210-0642	H1-BKG-2.5'-3'	20	91
-	Method Blank	0.10	N.D.

Analytes reported as N.D. were not present above the stated limit of detection.

**SEQUOIA ANALYTICAL** 

Pell

Tod Granicher Project Manager Please Note:

Analysis results reported on a dry-weight basis.

2100637.ENG <9>



Client Project ID: W.O. #4432

600 Bancroft Way Berkeley, CA 94710

Attention: Tom Paulson

QC Sample Group: 210-0637-42

Reported: Oct 20, 1992

## **QUALITY CONTROL DATA REPORT**

ANALYTE		Total Kjeldahl			
	Total Phosphorous	Nitrogen		<u> </u>	
Method: Analyst: Reporting Units: Date Analyzed: QC Sample #:	EPA365.3 K. Follett mg/kg Oct 19, 1992 210-0642	EPA351.4 G. Kern mg/kg Oct 15, 1992 210-0642			
Sample Conc.:	420	84	ing distribution of the second		· · · · · · · · · · · · · · · · · · ·
Spike Conc. Added:	100	4000			
Conc. Matrix Spike:	560	4000			
Matrix Spike % Recovery:	140	98			
Conc. Matrix Spike Dup.:	530	3800			
Matrix Spike Duplicate % Recovery:	110	93			
Relative % Difference:	5.6	5.1			

SEQUOIA ANALYTICAL

JUBAL

Tod Granicher Project Manager % Recovery:

Conc. of M.S. - Conc. of Sample x 100

Spike Conc. Added

Relative % Difference:

Conc. of M.S. - Conc. of M.S.D. x 100

(Conc. of M.S. + Conc. of M.S.D.) / 2

2100637.ENG <10>



600 Bancroft Way

Berkeley, CA 94710 Attention: Tom Paulson Client Project ID:

Lab Number:

W.O. #4432

210-0637

Sample Descript: Soil, H1-VW-4'-4.5'

Method of Analysis: ASTM D422-63

Sampled: Received:

Sep 2, 1992 Oct 7, 1992

Analyzed:

Oct 13, 1992

Reported:

Oct 20, 1992

## PARTICLE SIZE DISTRIBUTION BY SIEVE AND HYDROMETER

### SIEVE TEST

(A) TOTAL WEIGHT OF SAMPLE:

(B) WEIGHT RETAINED IN NO. 10 SIEVE:

(C) % PASSING NO. 10 SIEVE:

218.37g 33.38g 84.71

SIEVE TEST FOR WEIGHT RETAINED IN NO. 10 SIEVE

IDEAL PAN = 0.0 IDEAL TOTAL = (B)

	WEIGHT		CUMULATIVE	CUMULATIVE
SIEVE SIZE	RETAINED, g	% RETAINED	% RETAINED	% PASSING
1½ in.	0.0	0.0	0.0	100
3/8 in.	7.00	3.21	3.21	96.79
No. 4	10.54	4.83	8.04	91.96
No. 10	15.84	7.25	15.29	84.71
	i			
PAN	0.0	[	* * * * * * * * * * * * * * * * * * * *	
TOTAL	33.38	]		

#### HYDROMETER TEST

ELAPSED TIME	TEMP.	HYDROMETER	CORRECTED		PARTICLE
<b>(T)</b>	°C	READING (H)	READING (R)	(L)	DIAM. (S)
2	21	20	16	13.7	0.035
5	21	17	13	14.2	0.023
10	21	15	11	14.5	0.016
15	21	15	11	14.5	0.013
25	21	13	9	14.8	0.010
40	21	12	8	15.0	0.0083
60	21	11	7	15.2	0.0068
90	21.	11	7	15.2	0.0055
120	21	10	6	15.3	0.0048
1440	21	8	4	15.6	0.0014

% SUSPENDED
 (P) 21 17
21
17
15
15
12
11
9.3 9.3
9.3
8.0 5.3
5.3

WEIGHT OF SOIL USED IN HYDROMETER TEST (D): HYGROSCOPIC MOISTURE CORRECTION FACTOR (G): SPECIFIC GRAVITY (ASSUMED):

DISPERSING AGENT CORRECTION FACTOR (E):

MENISCUS CORRECTION FACTOR (F):

TEMP./SPEC. GRAVITY DEPENDANT CONSTANT (K):

	65g	l
į	0.983	l
	2.65	١
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	0.01348	İ

FORMULAS:

R = H - E - F

S = K[SQRT(L/T)]P = (R/W) 100

 $W = (J \cdot 100) / C$ 

 $J = D \cdot G$ 

SEQUOIA ANALYTICAL

Tod Granicher

Project Manager

2100637.ENG <1>

600 Bancroft Way

Client Project ID:

W.O. #4432

Sampled:

Sep 2, 1992

Berkeley, CA 94710

Sample Descript: Soil, H1-VW-4.5'-5.0' Method of Analysis: ASTM D422-63

Received: Analyzed:

Oct 7, 1992 Oct 13, 1992

Attention: Tom Paulson

Lab Number:

210-0638

Reported:

Oct 20, 1992

## PARTICLE SIZE DISTRIBUTION BY SIEVE AND HYDROMETER

#### SIEVE TEST

(A) TOTAL WEIGHT OF SAMPLE:

(B) WEIGHT RETAINED IN NO. 10 SIEVE:

(C) % PASSING NO. 10 SIEVE:

255.17g 4.75g 98.14

TOTAL

4.75

0.01348

SIEVE TEST FOR WEIGHT RETAINED IN NO. 10 SIEVE

> IDEAL PAN = 0.0 IDEALTOTAL = (B)

		WEIGHT		CUMULATIVE	CUMULATIVE
	SIEVE SIZE	· RETAINED, g	% RETAINED	% RETAINED	* % PASSING
I	1½ in.	0.0	0.0	0.0	100
-	3/8 in.	0.0	0.0	0.0	100
	No. 4	1.48	0.58	0.58	90.42
	No. 10	3.27	1.28	1.86	98.14
				l ·	
•	PAN	0.0		<u> </u>	

#### HYDROMETER TEST

ELAPSED TIME	TEMP.	HYDROMETER	CORRECTED		PARTICLE
(T)	°C	READING (H)	READING (R)	(L)	DIAM. (S)
2	21	12	8	15.0	0.037
5	21	10	6	15.3	0.024
10	21	9	5	15.5	0,017
15	21	8	4	15.6	0.014
25	21	8	4	15.6	0.011
40	21	. 8	4	15.6	0.0084
60	21	7	3	15.8	0.0069
. 90	21	7	3	15.8	0.0056
120	21	7	3	15.8	0.0049
1440	21	7	3	15.8	0.0014

% SUSPENDED 12 9.2 7.6 6.1 6.1 6.1 4.6 4.6 4.6 4.6

WEIGHT OF SOIL USED IN HYDROMETER TEST (D): HYGROSCOPIC MOISTURE CORRECTION FACTOR (G): SPECIFIC GRAVITY (ASSUMED):

DISPERSING AGENT CORRECTION FACTOR (E): MENISCUS CORRECTION FACTOR (F):

TEMP./SPEC. GRAVITY DEPENDANT CONSTANT (K):

65g FORMULAS: 0.983 R = H - E - FS = K[SQRT(L/T)]2.65 3 P = (R/W) 100 $W = (J \cdot 100) / C$ 1

 $J = D \cdot G$ 

**SEQUOIA ANALYTICAL** 

فالحظت



600 Bancroft Way

Berkeley, CA 94710 Attention: Tom Paulson Client Project ID:

W.O. #4432

Sample Descript: Soil, H1-A-3'-4'

Method of Analysis: ASTM D422-63 Lab Number: 210-0639

Sampled: Received: Sep 2, 1992 Oct 7, 1992

Analyzed: Reported:

Oct 13, 1992 Oct 20, 1992

## PARTICLE SIZE DISTRIBUTION BY SIEVE AND HYDROMETER

### SIEVE TEST

(A) TOTAL WEIGHT OF SAMPLE:

(B) WEIGHT RETAINED IN NO. 10 SIEVE:

(C) % PASSING NO. 10 SIEVE:

256.56a 99.81g

SIEVE TEST FOR WEIGHT RETAINED IN NO. 10 SIEVE

IDEAL PAN = 0.0 IDEALTOTAL = (B)

	WEIGHT		CUMULATIVE	CUMULATIVE
SIEVE SIZE	RETAINED, g	% RETAINED	% RETAINLD	% PASSING
11/2 in.	0.0	0.0	0.0	100
3/8 in.	41.21	16.06	16.06	83.94
No. 4	26.13	10.18 ***	26.24	73.76
No. 10	32.47	12.66	36.90	61.10
PAN	0.0	<u> </u>	<u> </u>	

TOTAL 99.81

#### HYDROMETER TEST

_ ELAPSED TIME	TEMP.	HYDROMETER	CORRECTED	•	PARTICLE
(T)	°C	READING (H)	READING (R)	(L)	DIAM. (S)
- 2	21	14	10	14.7	0.037
5	· 21	12	8	15.0	0.023
10	21	11	7	15.2	0.017
15	21	10	6	15.3	0.014
25	-21	9	5	15.5	0.011
40	21	9	5	15.5	0.0084
60	21	7	3	15.8	0.0069
90	21	7	3	15.8	0.0056
120	21	7	3	15.8	0.0049
1440	21	7	3	15.8	0.0014

% SUSPENDED 9.4 7.6 6.5 5.7 4.7 4.7 2.8 2.8 2.8 2.8

WEIGHT OF SOIL USED IN HYDROMETER TEST (D): HYGROSCOPIC MOISTURE CORRECTION FACTOR (G): SPECIFIC GRAVITY (ASSUMED):

DISPERSING AGENT CORRECTION FACTOR (E): MENISCUS CORRECTION FACTOR (F):

TEMP./SPEC. GRAVITY DEPENDANT CONSTANT (K):

65g	FORMULAS:
0.995	R=H-E-F
2.65	S = K[SQRT(L/T)]
3	P = (R/W) 100'
1	$W = (J \cdot 100) / C$
0.01249	1 1 - 10.00 ''

SEQUOIA ANALYTICAL



600 Bancroft Way

Client Project ID:

W.O. #4432

Sampled:

Sep 4, 1992

Berkeley, CA 94710

Sample Descript: Soil, H1-BKG-2.5'-3' Method of Analysis: ASTM D422-63

Received: Analyzed:

Oct 7, 1992 Oct 14, 1992

Attention: Tom Paulson

Lab Number:

210-0642

Reported:

Oct 20, 1992

## PARTICLE SIZE DISTRIBUTION BY SIEVE AND HYDROMETER

#### SIEVE TEST

(A) TOTAL WEIGHT OF SAMPLE:

(B) WEIGHT RETAINED IN NO. 10 SIEVE:

(C) % PASSING NO. 10 SIEVE:

280.69g 39.34g 85.98

TOTAL

SIEVE TEST FOR WEIGHT RETAINED IN NO. 10 SIEVE

IDEAL PAN = 0.0 IDEAL TOTAL = (B)

•	WEIGHT	•	CUMULATIVE	CUMULATIVE
SIEVE SIZE	RETAINED, g	% RETAINED	% RETAINED	% PASSING
1½ in.	0.0	0.0	0.0	100
.3/8 in.	24.76	8.62	8.82	91.18
No. 4	7.50	2.67	11.49	88.51
No. 10	7.08	2.52	14.01	85.99
		• •		
PAN	0.0			

#### HYDROMETER TEST

ELAPSED TIME	TEMP.	HYDROMETER	CORRECTED		PARTICLE
<u>(T)</u>	°C	READING (H)	READING (R)	(L)	DIAM. (S)
2	21	- 26	22	12.7	0.034
5	21	24	20	13.0	0.022
10	21	24	20	13.0	0.015
15	21	22	18	13.3	0.013
25	21	19	15	13.8	0.010
40	21	19	15	13.8	0.0079
60	21	15	11	14.5	0.0066
90	21	15	11	14.5	0.0054
120	21	12	8	15.5	0.0048
1440	21	9	5	15.5	0.0014

WEIGHT OF SOIL USED IN HYDROMETER TEST (D): HYGROSCOPIC MOISTURE CORRECTION FACTOR (G):

SPECIFIC GRAVITY (ASSUMED):

DISPERSING AGENT CORRECTION FACTOR (E):

MENISCUS CORRECTION FACTOR (F):

TEMP./SPEC. GRAVITY DEPENDANT CONSTANT (K):

65g	F
0.0984	ŀ
2.65	
3	
1	ĺ
0.01348	

39.34

FORMULAS:

R = H - E - FS = K[SQRT(L/T)]

P = (R/W) 100

 $W = (J \cdot 100) / C$ 

 $J = D \cdot G$ 

SEQUOIA ANALYTICAL

Tod Granicher

Project Manager

2100637.ENG <6>

600 Bancroft Way

Client Project ID:

W.O. #4432 Sample Descript: Soil, H2-VW-3'-3.5'

Sampled: Received: Sep 3, 1992

Berkeley, CA 94710

Method of Analysis: ASTM D422-63

Analyzed:

Oct 7, 1992 Oct 13, 1992

Attention: Tom Paulson

Lab Number:

210-0640

Reported:

Oct 20, 1992

### PARTICLE SIZE DISTRIBUTION BY SIEVE AND HYDROMETER

### SIEVE TEST

(A) TOTAL WEIGHT OF SAMPLE:

(B) WEIGHT RETAINED IN NO. 10 SIEVE:

(C) % PASSING NO. 10 SIEVE:

203.97a 33.37g

88.64

SIEVE TEST FOR WEIGHT RETAINED IN NO. 10 SIEVE

IDEAL PAN = 0.0 IDEALTOTAL = (B)

	WEIGHT		CUMULATIVE	CUMULATIVE
SIEVE SIZE	RETAINED, g	% RETAINED	% RETAINED	% PASSING
1½ in.	0.0	. 0.0	0.0	100
3/8 in.	4.04	1.96	1.98	98.02
No. 4	15.8 i	7.75	9.73	90.27
No. 10	13.52	6.63	16.36	83.64
_				
13.5.1			<del> </del>	

PAN 0.0 **TOTAL** 33.37

#### HYDROMETER TEST

ELAPSED TIME	TEMP.	HYDROMETER	CORRECTED		PARTICLE
(T)	°C	READING (H)	READING (R)	(L)	DIAM. (S)
2	21	9	5	15.5	0.038
5	21	9	5	15.5	0.024
10	21	8	4	15.6	0.017
15	21	7	3	15.8	0.014
25	21	7	3	15.8	0.011
40	21	7	3	15.8	0.0085
60	21	6	2	16.0	0.0070
90	21	5	1	16.1	0.0057
120	21	5	1	16.1	0.0049
1440	21	5	1	16.1	0.0014

% SUSPENDED
(P)
6.5
G.5
5.2
3.9
3.9
3.9
2.6
1.3
1.3
1.3

WEIGHT OF SOIL USED IN HYDROMETER TEST (D): HYGROSCOPIC MOISTURE CORRECTION FACTOR (G): SPECIFIC GRAVITY (ASSUMED):

DISPERSING AGENT CORRECTION FACTOR (E): MENISCUS CORRECTION FACTOR (F):

TEMP./SPEC. GRAVITY DEPENDANT CONSTANT (K):

FORMULAS: 65g 0.983 2.65 3 1 0.01348

R = H - E - FS = K[SQRT(L/T)]P = (R/W) 100 $W = (J \cdot 100) / C$ 

 $J = D \cdot G$ 

**SEQUOIA ANALYTICAL** 



600 Bancroft Way

Berkeley, CA 94710 Attention: Tom Paulson Client Project ID: Sample Descript:

Lab Number:

W.O. #4432 Soil. H2-VW-4'-4.5

Sample Descript: Soil, H2-VW-4'-4.5' Method of Analysis: ASTM D422-63 Sampled: Received: Sep 3, 1992

Analyzed:

Oct 7, 1992 Oct 14, 1992

Reported:

Oct 20, 1992

## PARTICLE SIZE DISTRIBUTION BY SIEVE AND HYDROMETER

210-0641

#### SIEVE TEST

(A) TOTAL WEIGHT OF SAMPLE:

(B) WEIGHT RETAINED IN NO. 10 SIEVE:

(C) % PASSING NO. 10 SIEVE:

228.60g 7.04g 96.92

SIEVE TEST FOR WEIGHT RETAINED IN NO. 10 SIEVE

IDEAL PAN = 0.0 IDEAL TOTAL = (B)

: SIEVE SIZE	WEIGHT	% RETAINED	CUMULATIVE % RETAINED	CUMULATIVE '% PASSING
1½ in.	0.0	0.0	0.0	# FASSING 100
3/8 in.	0.0	0.0	0.0	100
No. 4	2.50g	1.09	1.09 ···	98.91
No. 10	4.54g	1.99	3.08	95.92
PAN	0.0			
TOTAL	7.04			

#### **HYDROMETER TEST**

	ELAPSED TIME	TEMP.	HYDROMETER	CORRECTED		PARTICLE
	<u>(T)</u>	°C	READING (H)	READING (R)	(L)	DIAM. (S)
	2	21	7	3	15.8	0.038
	5	21	7	3	15.8	0.024
	10	21	7	3	15.8	0.017
	15	21	6	2	16.0	0.014
	25	21	5	1	16.1	-0.011
	40	. 21	5	1	16.1	0.0086
	60	21	5	1	16.1	0.0070
L	90	21	5	1	16.1	0.0057
	120	21	5	1	16.1	0.0049
	1440	21	5	1	16.1	0.0014

	% SUSPENDED
	(P)
	4.5
	4.5
	4.5
	3.0
. !	1.5
ļ	1.5
ı	1.5
- {	1.5
	1.5
	1.5

o cuencinen

WEIGHT OF SOIL USED IN HYDROMETER TEST (D): HYGROSCOPIC MOISTURE CORRECTION FACTOR (G):

SPECIFIC GRAVITY (ASSUMED):

DISPERSING AGENT CORRECTION FACTOR (E):

MENISCUS CORRECTION FACTOR (F):

TEMP./SPEC. GRAVITY DEPENDANT CONSTANT (K):

65g	I
0.995	l
2.65	İ
3	
1	
0.01348	

FORMULAS:

R = H - E - F

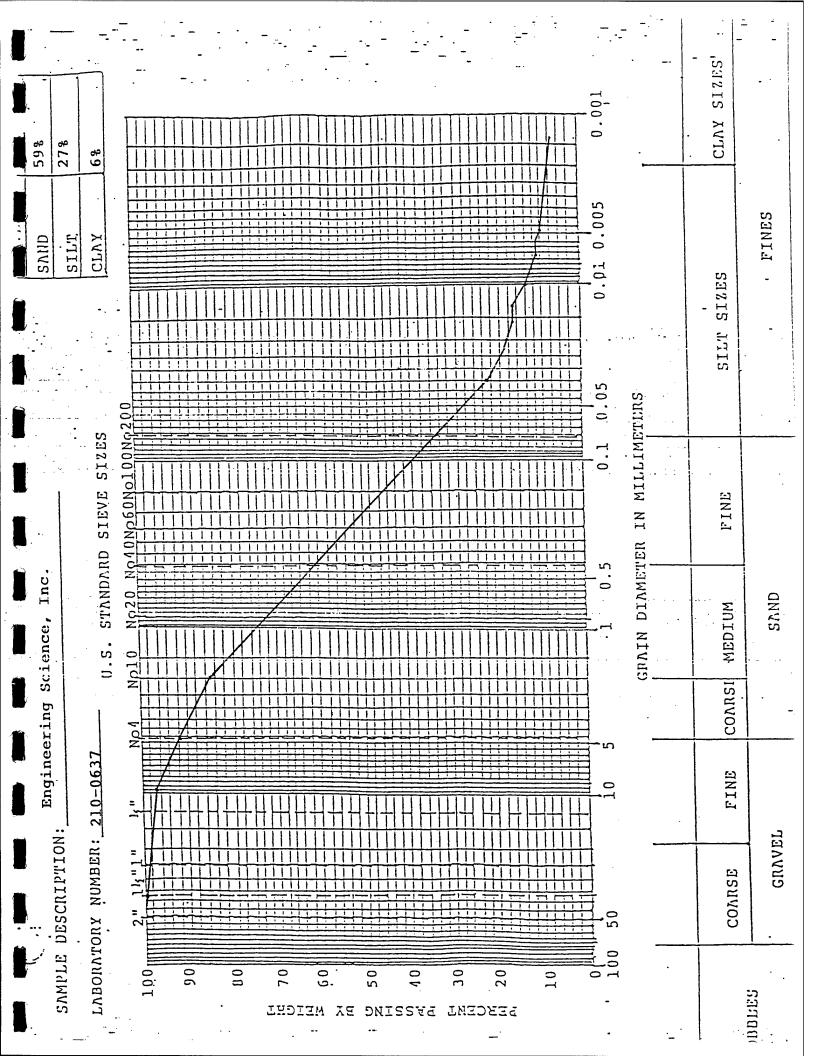
S = K[SQRT(L/T)] P = (R/W) 100

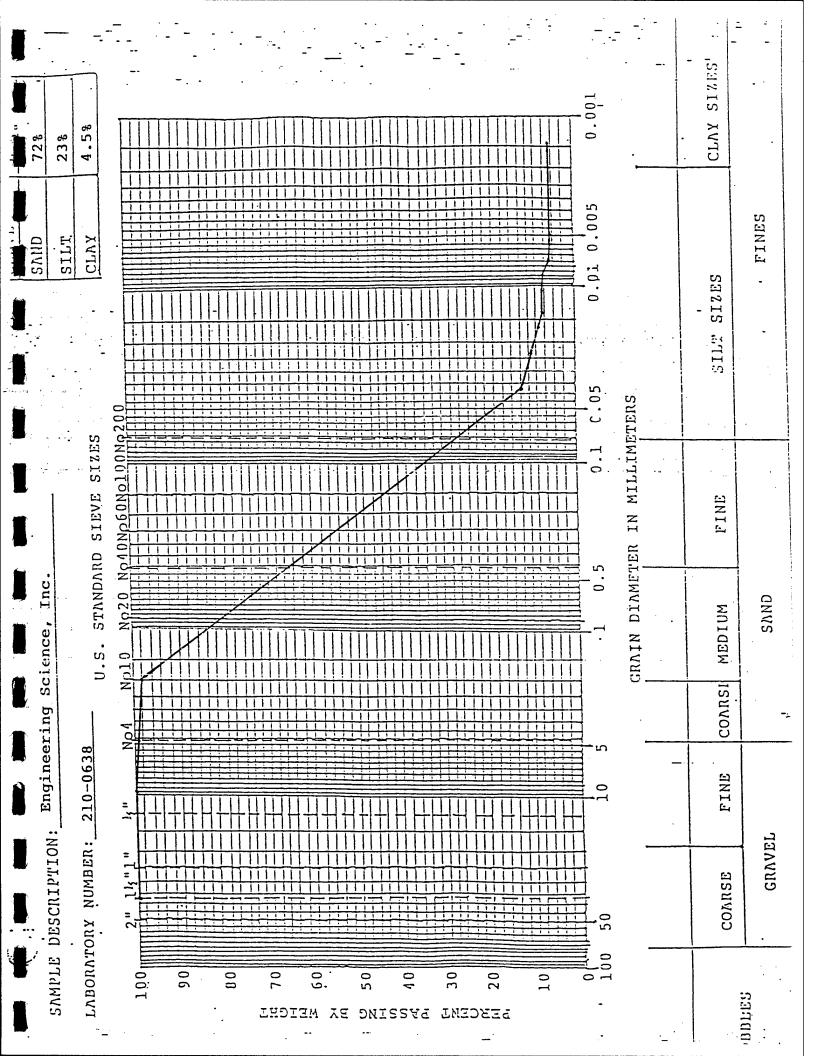
P = (R/W)100W = (J.100)/C

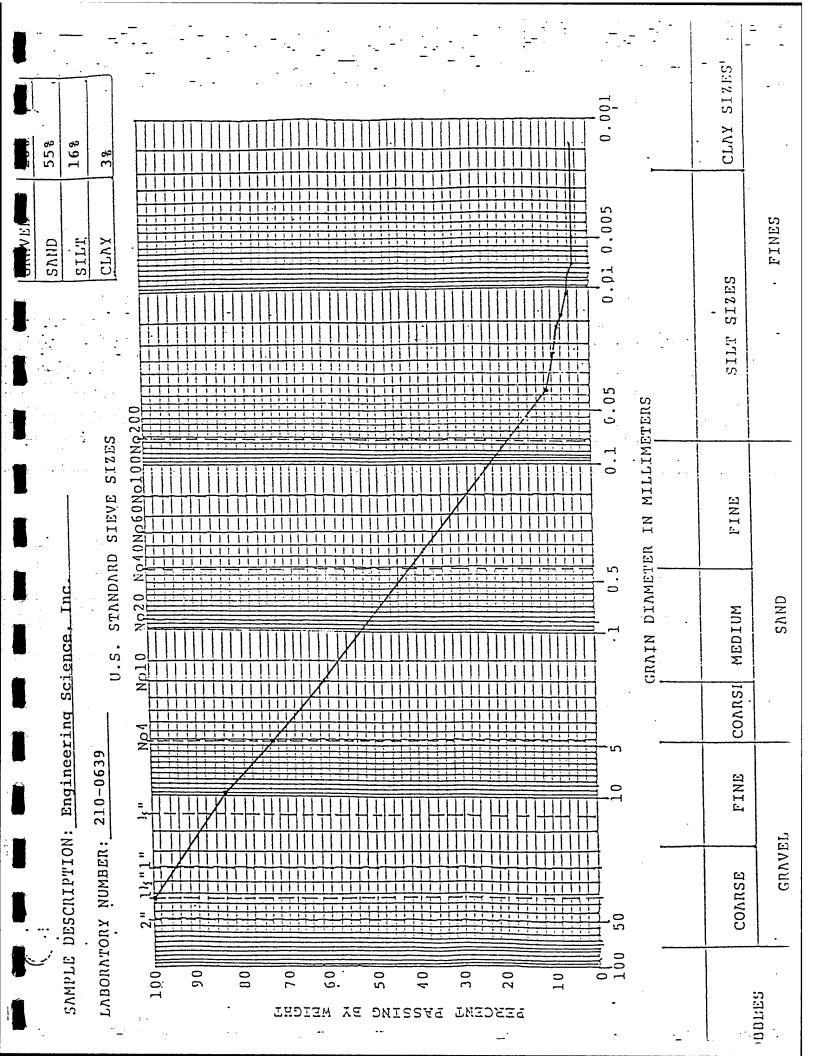
 $J = D \cdot G$ 

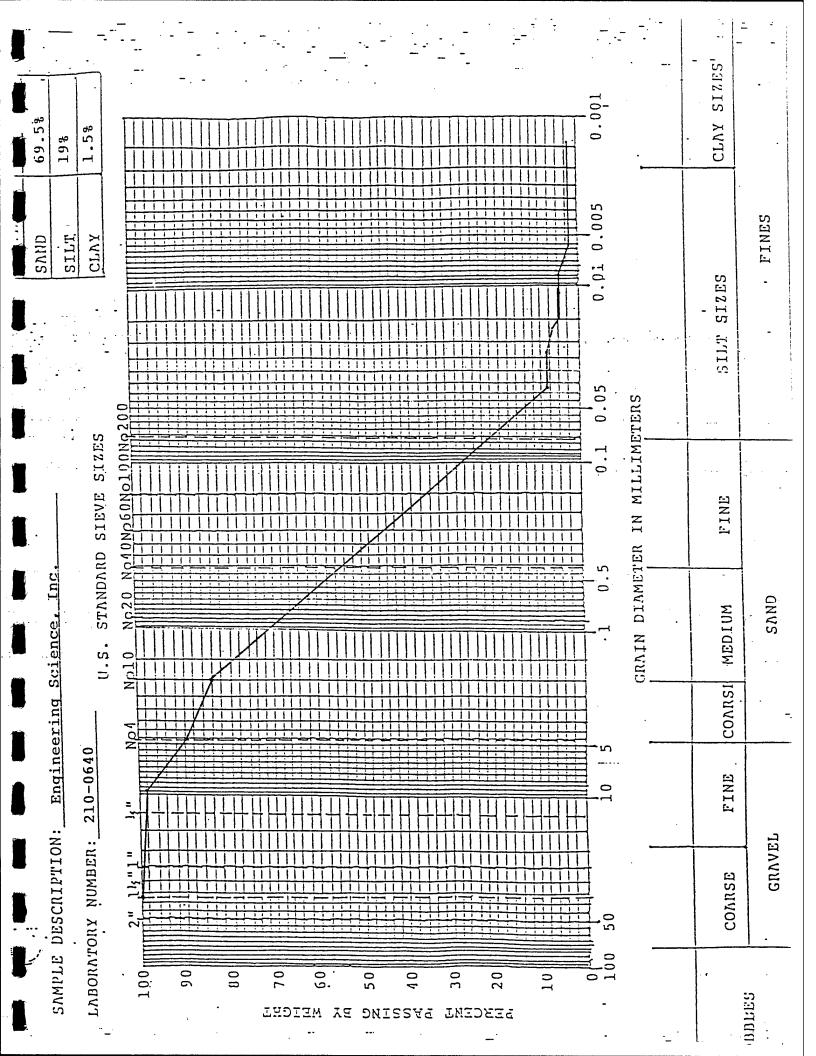
**SEQUOIA ANALYTICAL** 

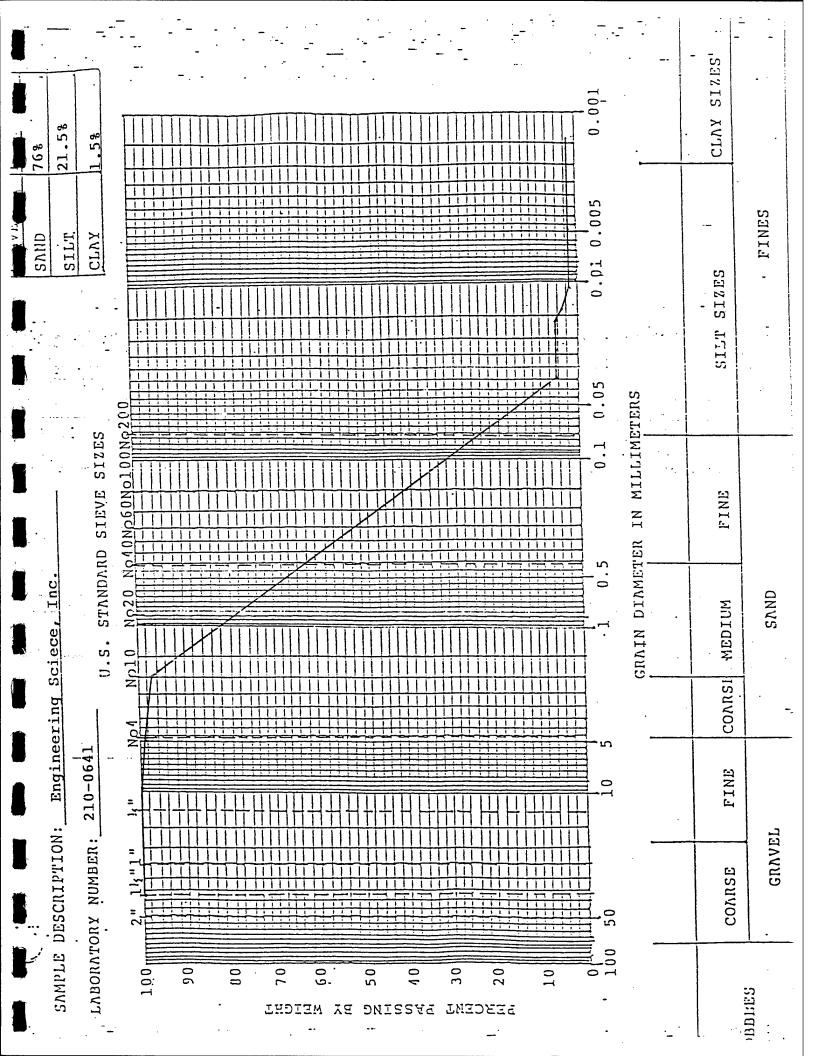
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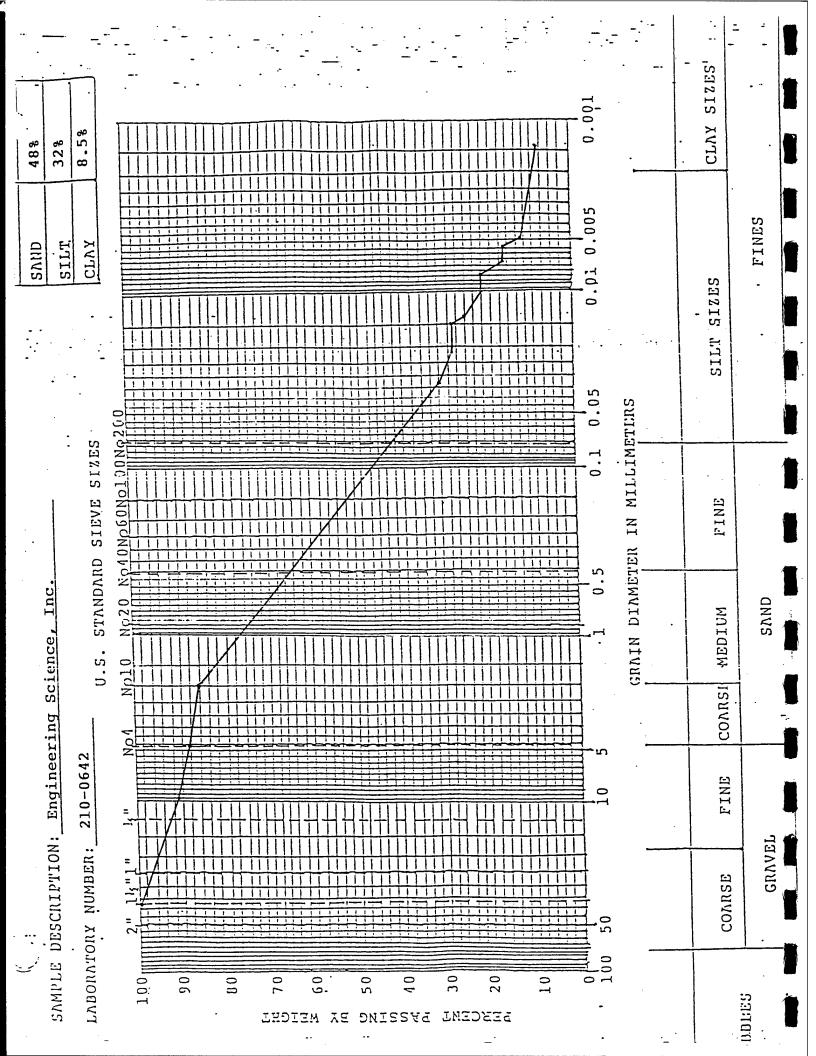












ENGINEERING-ECTENCE

CHAIN OF CUSTODY RECORD FOR WATER BAMPLES

											Τ
EB JOH	. 011	PROJECT NA	PROJECT NAME/LOCATION	•	PREB	PREBERVATIVES		REQUIRED	C		
		4432			5347	·     ·				To: STELLAN LAB.	
FIELD CC	CONTACT:					HNNLYEE	ев педи	кедил:кер		J	
BAHPLERB	в илиев	1 & BIGNATURES	33		0)51						<u>.</u>
			. •		47 31 15346		-				-
DATE	TIME	FIELD SAMPLE	E IDENTIFIER	7_	1315 74 <u>1</u>					пенликв	<del></del>
9/2/92 1	1560	HI-VW-4"	HI- VW- 4'-45' (4433,018)	1	7		001	633		Word nosult as Any	, STH
	1500	iti.vw-4.51-5	141.VW-4,51-5,0' (4432.028)	7.	<u>\</u>		Ŏ	XX		Exers. art MD/ 15. F.	12
'/	1.500	141-4-3-41	141-4-3-4' (4432038)		\		0			MajorTrug units.	
9/3/6,2 6	1200	H2-VW- 3'	42.VW- 3- 3.5 (4432,044)V	1	/			2		Urport METERS Blow	31
	*	42-VW-4-4.5	42-VW-4-45'(4422,058)	\	7		Ö	50	·	45/150 (10 day next	\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\\
4/62		H1-BKG- 2,	H1-BKG- 2,5-3 (443,068		7	ر— 	Ö	0)5		177.) Noport not a	16
										To Tay Paulson.	8
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"RECEIVED		FOR LABORATORY DX:		1	(	(		ZCI -	DATE: 10	16/92 TIME: 4:00 2m	T
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Battelle

JOB No. Engineering-Science Job No. VIE olubios

CHAIN OF CUSTC DY RECORD

Form No.

Columbus Laboratories					
	Project Title, Bio venting	SAMPLE TYPE (√)		1	
H 0+90-8747	HANSOM AFB	1 12/8	, ,		
SAMPLERS: (Signature)	15im Abott	Mital Lings of a	N 1anist	nadmuM fo fonistno	
DATE TIME	SAMPLE I.D.	MOST SENT	noo		Remarks
02 027 92	HI-VW - 4-45	メメメス		/	402 GIASS
02067 92	154-,h-M1-1H	* X X		/	1602, 61455
02 KT 92	H1-1W-4-4,5'	***			BRASS FUBE
03 ot 92	141-VW-4.5'-5.0'	<i>X X X X</i>		/	402 6/4SS
28 720EN	HI-WW-45-50	XXX		/	1602 6/455
22 CT 92	HI-VW-45'-5,0'	X		/	BUKS TUBE
CP 730 ES	HI-A-3-4'	× × × ×		/	402 GMSS
020CT 92	HI-A-3-4'	` \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \ \		, ,	160 6/HSS
26 170 00	H-A-38.31-4			/	BASS TUBE
030192	H2-VW-3'-3.5'			۲.	40 glass 160 6/185
130c7 92	H2-UW-3-3,51	X		/	SUKS TUBE
03057 92	13-1W-4/21	X X X X		/	402 6/ASS
h3 047 92	112-VW-4.5	×		-	1602 GIASS
03 OCT 92	H2-VW-4.5'	X		-	BRASS TUBE
24 KT 92	HI-BKG-2,5'-3'	XXX		_	402, GIASS
04 0CT 92	HI-BKG-2,5'-3'	$\times \times \times$			1602, 6/455
0400792	ווו				BRASS TUBE
Relinquished by: (Signature)	Date/Time Received by: (Signature)	ure) Relinquithed by: (Signature)	Date/Time	Received by:	;,
In Healy try	040x791 2200				
uishe	Date/Time Received by:	Refinquished by: (Signature)	Date/Time	Received by:	:^-
	(Signature)			₹ I	
Relinquished by: (Signature)	Date/Time Received for Laboratory by	Date/Time Ren	narks SEND RESOLTS JEFF K: HEL SOS KIMA AUE,	107 27	BATTELLE
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# APPENDIX C BUILDING 1639 SOIL GAS PERMEABILITY DATA

Table C-1. Results of Soil Gas Permeability Test at Monitoring Point H1-MPA

	Pressure ("H	Pressure ("H <sub>2</sub> O) by Depth		Pressure ("H	Pressure ("H <sub>2</sub> O) by Depth
Time (min)	2.5′	5.0′	Time (min)	2.5′	5.0′
0	0	0.01	20	0.025	0.32
₩-	0.05	0.17	23	0.025	0.33
2	0.02	0.21	26	0.025	0.34
33	0.02	0.22	29	0.03	0.35
4	0.03	0.225	32	0.025	0.35
5	0.033	0.235	35	0.03	0.35
9	0.35	0.25	38	0.025	0.35
7	0.03	0.27	41	0.025	0.35
∞	0.02	0.28	44	0.02	0.35
6	0.02	0.30	47	0.02	0.35
10	0.015	0.30	50	0.03	0.35
12	0.01	0.30	90	0.02	0.35
14	0.02	0.30	70	0.02	98:0
16	0.02	0.30	80	0.01	0.37
18	0.03	0.32	06	0.02	0.36

Table C-2. Results of Soil Gas Permeability Test at Monitoring Point H1-MPB

	Pressure ("H	Pressure ("H <sub>2</sub> O) by Depth		Pressure ("H	Pressure ("H <sub>2</sub> O) by Depth
Time (min)	2.5′	5.0′	Time (min)	2.5′	5.0′
1	0.045	0.01	12	0.080	0.015
2	90.0	0.02	50	0.115	0.025
3	0.065	0.01	55	0.115	0.025
4	0.07	0.015	09	0.110	0.02
5	0.075	0.02	70	0.115	0.02
9	0.075	0.02	08	0.113	0.02
7	0.090	0.015	06	0.110	0.022
8	0.092	0.015			
9	0.095	0.02			
10	0.085	0.02			

Table C-3. Results of Soil Gas Permeability Test at Monitoring Point H1-MPC

	Pressure ("H <sub>2</sub> O) by Depth	O) by Depth		Pressure ("H,O) by Depth	O) by Depth
Time (min)	3.5′	6.0′	Time (min)	3.5′	6.0′
0 .	0	0	20	0.01	0.02
<b>,</b>	0	0.01	23	0.02	0.015
2	0.01	0.015	26	0.03	0.035
3	0.01	0.03	29	0.03	0.015
4	0.02	0.02	32	0.02	0.02
5	0.03	0.025	35	0.025	0.015
9	0.02	0.03	38	0.015	0.02
7	0.03	0.01	41	0.025	0.15
&	0.01	0.015	44	0.01	0.01
6	0.015	0.005	47	0.005	0.015
10	0.01	0.02	50	0.02	0.04
12	0	0	09	0.01	0.02
14	0.02	0.01	70	0.01	0.015
16	0.015	0.025	80	0.01	0.01
18	0.03	0.02	09	0.015	0.01

# APPENDIX D BUILDING 1639 IN SITU RESPIRATION TEST DATA

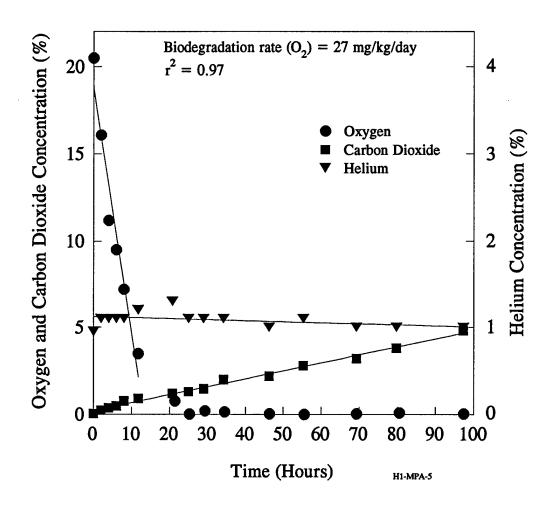


Figure D-1. Oxygen Utilization and Carbon Dioxide Production During the In Situ Respiration Test at Monitoring Point H1-MPA-5.0'

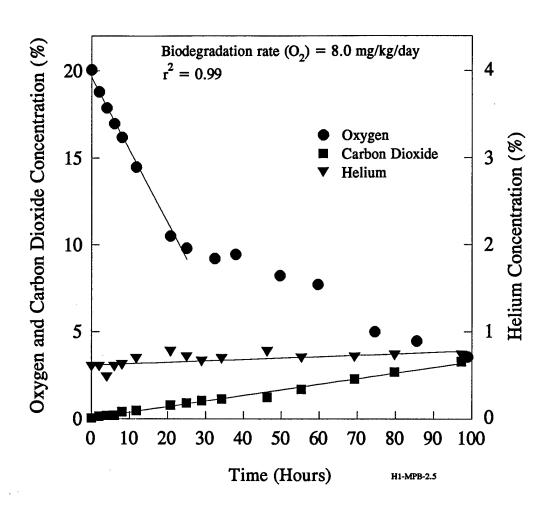


Figure D-2. Oxygen Utilization and Carbon Dioxide Production During the In Situ Respiration Test at Monitoring Point H1-MPB-2.5'

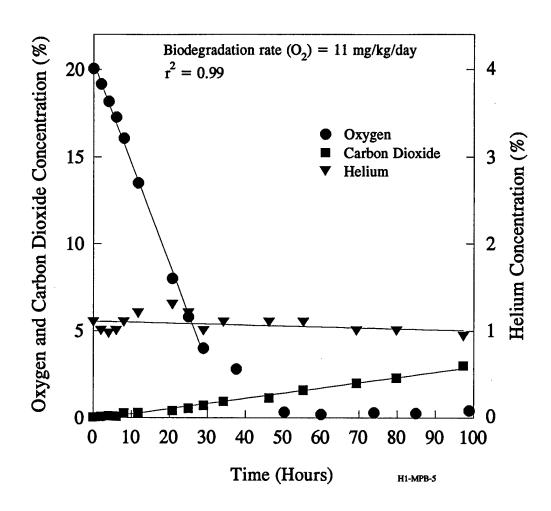


Figure D-3. Oxygen Utilization and Carbon Dioxide Production During the In Situ Respiration Test at Monitoring Point H1-MPB-5.0'

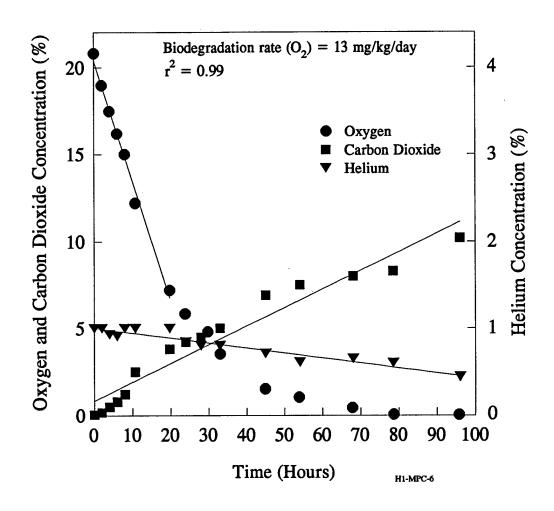


Figure D-4. Oxygen Utilization and Carbon Dioxide Production During the In Situ Respiration Test at Monitoring Point H1-MPC-6.0'

# APPENDIX E BUILDING 1812 SOIL GAS PERMEABILITY DATA

Table E-1. Results of Soil Gas Permeability Test at Monitoring Point H2-MPA

	Pressure ("H <sub>2</sub> O) by Depth	O) by Depth		Pressure ("H <sub>2</sub> O) by Depth	O) by Depth
Time (min)	2.5′	5.0′	Time (min)	2.5′	5.0′
0	0	0	20	0.55	0
2	0.50	0	25	0.55	0
3	0.54	0	30	0.56	0
4	0.55	0.005	35	0.56	0
5	0.55	0	40	0.56	0
9	0.55	0	95	0.57	0
7	0.55	0	09	0.58	0
8	0.55	0	75	0.58	0
6	0.55	0	06	0.59	0
12	0.55	0			
16	0.55	0.55			
18	0.55	0			

Table E-2. Results of Soil Gas Permeability Test at Monitoring Point H2-MPB<sup>1</sup>

	Pressure ("H	re ("H <sub>2</sub> O) by Depth		Pressure ("H <sub>2</sub> O) by Depth	O) by Depth
Time (min)	2.5′	5.0′	Time (min)	2.5′	5.0′
0	0	0	14	0.01	0.01
1	0	0	16	0	200'0
2	0	0	18	0.005	9000
3	0	0.005	20	0	0
4	0	0	25	0	0
5	0	0	30	0.005	200'0
9	0	0	35	0	0
7	0	0	40	0.005	500.0
8	0	0	20	0.005	9000
6	0	0	09	0.005	9000
10	0	0	75	0.005	9000
12	0.005	0.005			

Pressure readings were not collected from monitoring point H1-MPB-7.0'.

Table E-3. Results of Soil Gas Permeability Test at Monitoring Point H2-MPC

	Press	Pressure ("H <sub>1</sub> O) by Depth	Depth		Pressi	Pressure ("H <sub>2</sub> O) by Depth	Depth
Time (min)	2.5′	4.5′	6.0′	Time (min)	2.5'	4.5′	6.0′
1	0	0>	0>	16	0	0>	0>
2	0>	<0>	0	18	0	0>	0>
3	0>	0>	<0	20	0	0>	0>
4	<0>	<0>	<0	25	0	0>	0>
5	0>	0>	<0	30	0	0	0
9	0	0>	<0	35	0	0	0
7	0	<0>	<0	40	0	0	0
<b>∞</b>	0	0>	<0	20	0	0	0
6	0	<0>	<0	09	0	0	0
10	0	0>	<0	02	0	0	0
12	0	0>	<0	08	0	0	0
14	0	0>	<0>	06	0	0	0